

GenCore version 5.1.9
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On nucleic - nucleic search, using sw model

Run on: July 17, 2006, 20:07:43 ; Search time 2712 Seconds

(without alignments)
9431.773 Million cell updates/sec

Title: SEQ1-47502C

Perfect score: 399.6

Sequence: 1 ccaggtactcagcagcatgtgc.....tatgcagagagaccacaaag 400

Scoring table: IDENTITY_NUC

Gapop 10.0 , Gapext 1.0

Searched: 6366136 seqs, 31973710525 residues

Total number of hits satisfying chosen parameters: 12732272

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 150 summaries

Database :

GenEmbl: *
1: gb_env:*
2: gb_pat:*
3: gb_ph:*
4: gb_pl:*
5: gb_pr:*
6: gb_ro:*
7: gb_sts:*
8: gb_sy:*
9: gb_un:*
10: gb_vl:*
11: gb_ov:*
12: gb_hcg:*
13: gb_in:*
14: gb_cm:*
15: gb_da:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	398	99.6	187064	5	AC011476 Homo sapi
2	231	57.8	204340	12	AC019238 Homo sapi
3	161.2	40.3	184822	12	AC146499 Actus nan
4	160.6	40.2	232406	12	AC146898 Pongo pyg
5	160	40.0	97676	5	AC005497 Homo sapi
6	159.8	40.0	107866	5	AL445686 Human DNA
7	159.4	40.0	171480	5	AC016542 Homo sapi
8	159.4	39.9	97916	5	Z98048 Human DNA
9	158.8	39.7	195616	12	AC090344 Homo sapi
10	158.8	39.7	197156	12	AC090345 Homo sapi
11	158.8	39.7	197856	12	AP000777 Homo sapi
12	158.6	39.7	68990	12	AC087662 Homo sapi
13	158.4	39.6	96593	2	DD164421 NOVEL.COM
14	158.4	39.6	96593	2	AX695641 Sequence
15	158.4	39.6	112659	5	AC010677 Homo sapi
16	158.2	39.6	155344	5	AC026407 Homo sapi
17	157.8	39.5	170154	5	AC040168 Homo sapi
18	157.8	39.5	186747	12	AC018605 Homo sapi

19	157.8	39.5	198575	5	AC007495 Homo sapi
20	157.8	39.5	224187	5	AL732374 Human DNA
21	157.6	39.4	163908	12	AC150910 Pan trogl
22	157.4	39.4	174097	5	AC069513 Homo sapi
23	157.4	39.3	209844	5	AC011495 Homo sapi
24	157.2	39.3	173556	12	AC074388 Homo sapi
25	157	39.3	84001	5	AL162739 Human DNA
26	156.8	39.2	158330	5	AC025599 Homo sapi
27	156.8	39.2	195070	12	AC068995 Homo sapi
28	156.6	39.2	111372	12	AL353145 Homo sapi
29	156.6	39.2	150934	12	AC022252 Homo sapi
30	156.6	39.2	163521	12	AC108040 Homo sapi
31	156.6	39.2	196573	5	AC114480 Homo sapi
32	156.4	39.1	102008	12	AC016552 Homo sapi
33	156.4	39.1	244525	5	AC008499 Homo sapi
34	156.4	39.0	88848	5	AC107377 Homo sapi
35	155.8	39.0	182725	12	AC172796 Homo sapi
36	155.8	39.0	216387	5	DJ534K4
37	155.6	38.9	224187	5	AL732374 Homo sapi
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39	155.4	38.9	126462	5	AC004876 Homo sapi
40	155.4	38.9	132492	5	AC007616 Homo sapi
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42	155.4	38.9	183444	5	AP001024 Homo sapi
43	155.4	38.9	202138	12	AC018423 Homo sapi
44	155.4	38.9	204493	5	AC099489 Homo sapi
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48	155.2	38.8	178079	12	AC068929 Homo sapi
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50	155	38.8	72433	5	AC068889 Homo sapi
51	155	38.8	166884	12	AC023985 Homo sapi
52	155	38.8	178111	5	AC011739 Homo sapi
53	155	38.8	196216	5	AC099343 Homo sapi
54	154.8	38.7	98832	5	AL137127 Homo sapi
55	154.8	38.7	152757	5	AC018475 Homo sapi
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58	154.4	38.6	159273	12	AC025001 Homo sapi
59	154.4	38.6	169714	12	AC138876 Homo sapi
60	154.4	38.6	182892	5	AC034244 Homo sapi
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64	154.2	38.6	122748	5	AL031846 Human DNA
65	154.2	38.6	161166	12	AC087673 Homo sapi
66	154	38.5	20425	12	BX284653_5
67	154	38.5	110000	12	BX284653_4
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71	154	38.5	176813	12	AC107895 Homo sapi
72	154	38.5	198821	5	AC120057 Homo sapi
73	154	38.5	206773	12	AC010932 Homo sapi
74	153.8	38.5	115246	12	AC161015 Homo sapi
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76	153.8	38.5	154616	5	AC004846 Homo sapi
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79	153.8	38.5	203046	12	AC006342 Homo sapi
80	153.8	38.5	206258	12	AC112775 Homo sapi
81	153.8	38.5	233418	12	AC109128 Homo sapi
82	153.6	38.4	139063	12	AC133553 Homo sapi
83	153.6	38.4	146208	5	AL354776 Human DNA
84	153.6	38.4	167101	5	AC002094 Homo sapi
85	153.6	38.4	174724	12	AC140096 Pan trogl
86	153.6	38.4	176181	5	AC008155 Homo sapi
87	153.6	38.4	184684	12	AC148258 Homo sapi
88	153.4	38.4	67396	5	AC008848 Homo sapi
89	153.4	38.4	141605	5	AL355353 Human DNA
90	153.4	38.4	161251	12	AC147284 Pan trogl
91	153.4	38.4	166697	12	AC021103 Homo sapi

AC007495 Homo sapi	AL732374 Human DNA
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AC087673 Homo sapi	Continuation (6 of 6)
Continuation (5 of 6)	AL450328 Human DNA
AL365504 Human DNA	AC026834 Homo sapi
AC107895 Homo sapi	AC120057 Homo sapi
AC010932 Homo sapi	AC161015 Homo sapi
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AL355353 Human DNA	AC147284 Pan trogl
AC021103 Homo sapi	

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C	97	153.2	38.3	183411	12	AC137495	AC137495 Homo sapi
C	98	153.2	38.3	184453	5	AC025593	AC025593 Homo sapi
C	99	153.2	38.3	270567	12	AC138086	AC138086 Homo sapi
C	100	153	38.3	34335	5	AC112711	AC112711 Homo sapi
C	101	153	38.3	170758	5	AC004965	AC004965 Homo sapi
C	102	152.8	38.2	102562	12	AC109516	AC109516 Homo sapi
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C	104	152.8	38.2	166942	5	HS1054A22	AL031651 Human DNA
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C	110	152.6	38.2	146741	5	AC011452	AC011452 Homo sapi
C	111	152.6	38.2	168931	12	AC144998	AC144998 Pan trogl
C	112	152.6	38.2	169074	12	AC148830	AC148830 Pan trogl
C	113	152.6	38.2	175999	12	AC021154	AC021154 Homo sapi
C	114	152.6	38.2	190430	12	AC118987	AC118987 Pan trogl
C	115	152.4	38.1	576	7	BV198237	BV198237 sqm19689
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C	118	152.4	38.1	142170	5	AC087362	AC087362 Homo sapi
C	119	152.4	38.1	143847	5	AF205589	AF205589 Homo sapi
C	120	152.4	38.1	160864	12	AC021993	AC021993 Homo sapi
C	121	152.4	38.1	162835	5	AC087623	AC087623 Homo sapi
C	122	152.4	38.1	163494	5	AC079865	AC079865 Homo sapi
C	123	152.4	38.1	166237	5	AL139379	AL139379 Human DNA
C	124	152.4	38.1	170338	12	AF214633	AF214633 Homo sapi
C	125	152.4	38.1	174562	12	AC005848	AC005848 Homo sapi
C	126	152.4	38.1	194646	12	AC090129	AC090129 Homo sapi
C	127	152.4	38.1	205577	12	AC011237	AC011237 Homo sapi
C	128	152.2	38.1	615	7	BV504898	BV504898 pqd33e03
C	129	152.2	38.1	51242	5	BX120002	BX120002 Human DNA
C	130	152.2	38.1	149481	5	AC013472	AC013472 Homo sapi
C	131	152.2	38.1	155735	5	AC007683	AC007683 Homo sapi
C	132	152.2	38.1	159726	5	AC125257	AC125257 Homo sapi
C	133	152.2	38.1	168017	5	BS000238	BS000238 Pan trogl
C	134	152.2	38.1	169604	5	AC083884	AC083884 Homo sapi
C	135	152.2	38.1	175303	12	AC024721	AC024721 Homo sapi
C	136	152.2	38.1	178345	5	AC091172	AC091172 Homo sapi
C	137	152.2	38.1	190477	5	PTB109N16	AL954242 Pan trogl
C	138	152.2	38.1	196735	12	AC012192	AC012192 Homo sapi
C	139	152.2	38.1	197503	12	AC165190	AC165190 Colobus g
C	140	152	38.0	147118	5	AC007513	AC007513 Homo sapi
C	141	152	38.0	174424	2	AX335950	AX335950 Sequence
C	142	152	38.0	179676	12	AC129889	AC129889 Papio anu
C	143	152	38.0	181343	5	US2112	US2112 Homo sapien
C	144	152	38.0	184899	12	AC113392	AC113392 Homo sapi
C	145	152	38.0	223606	12	AC013479	AC013479 Homo sapi
C	146	152	38.0	225370	5	AL445222	AL445222 Human DNA
C	147	151.8	38.0	169323	12	AL161906	AL161906 Homo sapi
C	148	151.8	38.0	169515	5	AP002777	AP002777 Homo sapi
C	149	151.6	37.9	743	7	BV642952	BV642952 S216P6376
C	150	151.6	37.9	40536	12	AC110593	AC110593 Homo sapi

ALIGNMENTS

RESULT 1
LOCUS AC011476 187064 bp DNA linear PRI 16-AUG-2002
DEFINITION Homo sapiens chromosome 19 clone CTC-550B14, complete sequence.
ACCESSION AC011476
VERSION AC011476.8 GI:22267569
KEYWORDS
SOURCE HTG.
ORGANISM Homo sapiens (human)

REFERENCE	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini; Homiidae; Homo.
AUTHORS	1 (bases 1 to 187064)
JOURNAL	DOE Joint Genome Institute and Stanford Human Genome Center.
TITLE	Direct Submission
REFERENCE	Unpublished
AUTHORS	2 (bases 1 to 187064)
JOURNAL	DOE Joint Genome Institute.
TITLE	Direct Submission
REFERENCE	Submitted (07-OCT-1999) Production Sequencing Facility, DOE Joint Genome Institute, 2800 Mitchell Drive, Walnut Creek, CA 94598, USA
AUTHORS	3 (bases 1 to 187064)
JOURNAL	DOE Joint Genome Institute and Stanford Human Genome Center.
TITLE	Direct Submission
REFERENCE	Submitted (16-AUG-2002) DOE Joint Genome Institute, 2800 Mitchell Drive, Walnut Creek, CA 94598, USA
AUTHORS	On Aug 16, 2002 this sequence version replaced gi:15022008.
JOURNAL	Draft Sequence Produced by DOE Joint Genome Institute
COMMENT	www.jgi.doe.gov Finishing Completed at Stanford Human Genome Center www.sngc.stanford.edu Quality: Phrap Quality >=40 99.9% of Sequence; Estimated Total Number of Errors is 0.3. NOTE: Transposon sequencing failed to verify the number of repeat copies 67315-69359. Unsure number of repeat copies 67315-69359. NOTE: Shatter libraries failed to resolve dinucleotide repeat region 171590-171722. Unsure number of repeat copies 171590-171722. Forced join 171695.
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QY	1 CCAGTACTCGACGATGCTGGGCGATGGGACCAATTTATATAGATTGTACG 60
DB	119458 CCAGTACTCGACGATGCTGGGCGATGGGACCAATTTATATAGATTGTACG 119517
QY	61 CCAGGATGACACTGCTGTAATGCTGTATATCCAGACTTTCGGAGGCGCAAGTGGCG 120
DB	119518 CCAAGCATGACACTGCTGTAATGCTGTATATCCAGACTTTCGGAGGCGCAAGTGGCG 119577
QY	121 GATCACTGAGGTCAAGATCGAGACCATCTTGGCCCAACTGTGTAACCCCGCTTTA 180
DB	119578 GATCACTGAGGTCAAGATCGAGACCATCTTGGCCCAACTGTGTAACCCCGCTTTA 119637
QY	181 CTAATAATACAAAATAATAGCTGGGCGATGGGACACACCTGTAGTCCAGTACTACG 240
DB	119638 CTAATAATACAAAATAATAGCTGGGCGATGGGACACACCTGTAGTCCAGTACTACG 119697
QY	241 AGCCGAGATTCAGTGAAGTGAATCGCAGAGTGGCCGAATATCAGATCAGAGTGG 300
DB	119698 AGCCGAGATTCAGTGAAGTGAATCGCAGAGTGGCCGAATATCAGATCAGAGTGG 119757
QY	301 AGCAGATGAGACCCGCTCTCAAAAACAACAACAAAAAACCAATTAAGCATTTG 360
DB	119758 AGCAGATGAGACCCGCTCTCAAAAACAACAACAAAAAACCAATTAAGCATTTG 119817

Oy 361 TCACATCGGCTTCCAGACTATTGACGAGACCAAAAAG 400
 DB 119818 TCACATCGGCTTCCAGACTATTGACGAGACCAAAAAG 119857
 RESULT 2
 AC019238
 LOCUS
 DEFINITION Homo sapiens chromosome 19 clone RP11-700B5, WORKING DRAFT
 AC019238 204340 bp DNA linear HTG 17-AUG-2000
 AC019238
 AC019238
 AC019238.5 GI:9838316
 HTG; HTGS PHASE1; HTGS_DRAFT.
 SOURCE Homo sapiens (human)
 ORGANISM Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini;
 Homidae; Homo.
 1 (bases 1 to 204340)
 Watserson, R.H.
 The sequence of Homo sapiens clone
 Unpublished
 2 (bases 1 to 204340)
 Watserson, R.H.
 Direct Submission
 Submitted (30-DEC-1999) Genome Sequencing Center, Washington
 University School of Medicine, 4444 Forest Park Parkway, St. Louis,
 MO 63108, USA
 On Aug 17, 2000 this sequence version replaced gi.9280808.

----- Genome Center -----
 Center: Washington University Genome Sequencing Center
 Center code: MUGSC
 Web site: http://genome.wustl.edu/gsc/index.shtml
 Project information -----
 Center project name: H.NH0700805
 Summary Statistics -----
 Sequencing vector: M13; 84k
 Sequencing vector: plasmid; 16k
 Chemistry: Dye-terminator Big Dye; 16k of reads
 Chemistry: Dye-terminator Big Dye; 16k of reads
 Assembly program: Phrap; version 0.990319
 Consensus quality: 19676 bases at least Q40
 Consensus quality: 200928 bases at least Q30
 Consensus quality: 201749 bases at least Q20
 Insert size: 236000; agarose-fp
 Insert size: 204523; sum-of-contigs
 Quality coverage: 6.54 in Q20 bases; agarose-fp
 Quality coverage: 7.57 in Q20 bases; sum-of-contigs

 * NOTE: This is a 'working draft' sequence. It currently
 * consists of 12 contigs. The true order of the pieces
 * is not known and their order in this sequence record is
 * arbitrary. Gaps between the contigs are represented as
 * runs of N, but the exact sizes of the gaps are unknown.
 * This record will be updated with the finished sequence
 * as soon as it is available and the accession number will
 * be preserved.
 * 1 2752: contig of 2752 bp in length
 * 2753 2852: gap of unknown length
 * 2853 5901: contig of 3049 bp in length
 * 5902 6002: gap of unknown length
 * 6002 12965: contig of 6964 bp in length
 * 12966 13066: gap of unknown length
 * 13066 21018: contig of 7953 bp in length
 * 21019 21119: gap of unknown length
 * 21119 30399: contig of 9281 bp in length
 * 30400 30499: gap of unknown length
 * 30500 40958: contig of 10459 bp in length
 * 40959 41059: gap of unknown length
 * 41059 52180: contig of 11122 bp in length
 * 52181 52280: gap of unknown length

FEATURES
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 /clone="RP11-700B5"
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 clone end: SP6
 vector side: left
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 Best Local Similarity 99.1%; Pred. No. 8.9e-69;
 Matches 231; Conservative 1; Mismatches 1; Indels 0; Gaps 0;
 Oy 168 AACCCGCTTTACTAAATAATACAAAAATATGCTGGGAGTGTGACACACCTGTAGTC 227

Db		65376	AACCCGCTTTACTAATAAAATPACAATAAAATAGCTGGGCATGTGTGGCACAACCTGTAGTC	654359
Oy		228	CCAGCTACTCAGAGAGCCCGAGATTGCAGTGAAGTGGCAGAGTGAGCCGAATTCAC	287
Db		65436	CCAGCTACTCAGAGAGCCCGAGATTGCAGTGAAGTGGCAGAGTGAGCCGAATTCAC	654959
Oy		288	AGATCACAGAGTGGAGCAGAGTGAAGACACCCGCTCAAACAAACAACAAAAACAAAACAAA	347
Db		65496	AGATCACAGAGTGGAGCAGAGTGAAGACACCCGCTCAAACAAACAACAAAAACAAAACAAA	655555
Oy		348	CCATTAAGCATTTGCTCCATCTGCCGTTCCCAAGCTATTGTCAGAGACCAAAAAG	400
Db		65556	CCATTAAGCATTTGCTCCATCTGCCGTTCCCAAGCTATTGTCAGAGACCAAAAAG	65608
RESULT 3				
LOCUS	AC146499/c			
DEFINITION	Aotus nancymae clone CH258-450E24, WORKING DRAFT SEQUENCE, 2			
ACCESSION	AC146499	184822 bp	DNA	linear HTG 26-NOV-2003
VERSION	AC146499.1			
KEYWORDS	HTG; HTGS_PHASE2; HTGS_DRAFT.			
SOURCE	Aotus nancymae (Ma's night monkey)			
ORGANISM	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Euarchontoglires; Primates; Platyrrhini; Cebidae; Aotinae; Aotus.			
REFERENCE	1 (bases 1 to 184822)			
AUTHORS	Cheng,J.-F., Hamilton,M., Peng,Y., Mukherjee,S., Hosseini,R., Peng,Z., Malinov,I. and Rubin,E.M.			
TITLE	Direct Submission			
JOURNAL	2 (bases 1 to 184822)			
REFERENCE	Cheng,J.-F., Hamilton,M., Peng,Y., Mukherjee,S., Hosseini,R., Peng,Z., Malinov,I. and Rubin,E.M.			
AUTHORS	Direct Submission			
TITLE	Submitted (21-AUG-2003) Genome Sciences, Lawrence Berkeley National Laboratory, 1 Cyclotron Rd., Berkeley, CA 94720, USA			
JOURNAL	3 (bases 1 to 184822)			
REFERENCE	Cheng,J.-F., Hamilton,M., Peng,Y., Mukherjee,S., Hosseini,R., Peng,Z., Malinov,I. and Rubin,E.M.			
AUTHORS	Direct Submission			
TITLE	Submitted (26-NOV-2003) Genome Sciences, Lawrence Berkeley National Laboratory, 1 Cyclotron Rd., Berkeley, CA 94720, USA			
JOURNAL				
COMMENT				
	Sequence Produced by Berkeley PCA			
	Web site: http://pga.lbl.gov			
	Center Code: PGABERK			
	Center Project Name: W033			
	Bac Clone Name: CH258-450E24			
	<p>This sequence has been compared to sequences of other species using VISTA (http://www-gsd.lbl.gov/VISTA/). The results can be viewed at: http://pga.lbl.gov/cgi-bin/search_cvcgdr?pe=nk&value=ABOAL</p>			
	<p>The order-orientation of the draft sequence was accomplished by using: Avid (http://baboon.math.berkeley.edu/navid/), Lagan (http://lagan.stanford.edu/) and paired end information.</p>			
	Funding agent: Programs for Genomic Applications (NHLBI)			
	<p>Summary Statistics:</p> <p>Sequencing vector: plasmid; pUC18</p> <p>Chemistry: Dye-terminator Big Dye Assembly Program: Phrap version 0.990329.</p> <p>* NOTE: This is a 'working draft' sequence. It currently consists of 2 contigs. Gaps between the contigs are represented as runs of N. The order of the pieces is believed to be correct as given, however the sizes</p>			

	* If the gaps between them are based on estimates that have
	* provided by the submittor.
	* This sequence will be replaced
	* by the finished sequence as soon as it is available and
	* the accession number will be preserved.
*	1 32135: contig of 32135 bp in length
*	32136 32235: gap of unknown length
*	32236 184822: contig of 152587 bp in length.
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Best Local Similarity	75.9%; Pred. No. 4,7e-44;
Matches 211; Conservative	1; Mismatches 64; Indels 2; Gaps 1;
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DB	133786 AGCGTTGATGCCAGCACTTTGGAGAGCCAAAGCGCGTGATGC--GAGGTCAAAGAGA 133722
QY	141 TCAGACCAATCTCTGGCCAAATGTGTAAAACCCCGTCTTTACTTAATAATCAAAAAATTAGC 200
DB	133728 TCGAACCATCTGTGTCAACAATGGTGAANAACCGTCTTCACTTAATAATCAAAAAATTAGC 133665
QY	201 TGGGATGATGGTCACACACTGTAGTCCAGCTACTCAGAGCCGAGATTCAGTGTAGC 260
DB	133668 TGGGATGATGGTCACACACTGTATTCACACTCTTGGAGAGCTGATGACAGAGATTG 133605
QY	261 TGAGATCGCAGAGTAGAGCCGAATATCACAGATCACAGAGTAGAGAGACCCGCTT 320
DB	133608 CTGTAACCCAGAGAGCGGAGGTGTCCTCAGCTCTGGTAAACAAGATGAAACTGTCT 133545
QY	321 CAAAAACAACAACAAAAAACAAAAACCATTAAGCAT 358
DB	133548 CAAAAAAGAAAAAAGAAAAAGAAAGATAAATCTT 133511
RESULT 4	
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DEFINITION	Pongo pygmaeus clone CH253-50L14, WORKING DRAFT SEQUENCE, 4 ordered pieces.
ACCESSION	AC146898 GI:38638702
VERSION	HTG; HTGS PHASE2; HTGS DRAFT.
KEYWORDS	Pongo pygmaeus (orangutan)
SOURCE	Pongo pygmaeus
ORGANISM	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini; Homnidae; Pongo.
REFERENCE	1 (bases 1 to 232406)
AUTHORS	Antonellis,A., Ayele,K., Benjamin,B., Blakeley,R.W., Bouffard,G.G., Brinkley,C., Brooke,S., Chu,G., Coleman,B., Coleman,H., Engle,J., Granite,S., Guan,X., Gupta,J., Haghighi,P., Han,J., Hansen,N., Ho,S.-L., Hu,P., Hurle,B., Idol,J.R., Karlins,E., Kwong,P., Latic,P., Larson,S., Lee-Jin,S.-O., Legaspi,R., Maduro,Q.L., Maduro,V.B., Margulies,E.H., Massello,C., Maskeri,B., McDowell,J., Mullikin,J.C., Paguirigan,C., Pearson,R., Portnov,M.E., Prasad,A., Reddik-Dugue,N., Schandler,K., Schneider,M.G., Shah,K., Sison,C., Stancirpop,S., Thomas,J.W., Thomas,P.T., Tsipouri,V., Vogt,J.L., Wetherby,K.D., Young,A. and Green,E.D.
TITLE	NISC Comparative Sequencing Initiative
JOURNAL	Unpublished .
REFERENCE	2 (bases 1 to 232406)
AUTHORS	Green,E.D.
TITLE	Direct Submission


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QY 141 TCGAGACCATCTCTGGCCCAATGTGTAACCCCGCTTTACTTAAATAACAAAAATAGC 200
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DB 36193 TCAAGACCAAGCTGGCCCAATGTGTAACCCCGCTTTACTTAAATAACAAAAATAGC 36252
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DB 36253 CGGGCAATGTGGCAACATGCTGTATCCAGTACTCGGAGGCGTGAAGAGGAGACTG 36312
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RESULT 6
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 LOCUS Human DNA sequence from clone Rpl1-496D1 on chromosome 1 contains
 DEFINITION the gene for a novel protein (RPL26), a ribosomal protein L26
 (RPL26) pseudogene, the 5' end of the SRM1 gene for
 betine/arginine repetitive matrix 1 and a CpG island, complete
 sequence.
 AL445686
 ACCESSION AL445686.14 GI:12666298
 VERSION HTG: FLJ42528: RPL26: SRM1.
 KEYWORDS Homo sapiens (human)
 SOURCE Homo sapiens
 ORGANISM Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini;
 Homidae; Homo.
 REFERENCE 1 (bases 1 to 107668)
 AUTHORS Coville, G.
 TITLE Direct Submission
 JOURNAL Submitted (13-MAY-2005) Wellcome Trust Sanger Institute, Hinxton,
 Cambridgeshire, CB10 1SA, UK. E-mail enquiries: vegas@sanger.ac.uk
 On Feb 5, 2001 this sequence version replaced gi:12581050.
 The following abbreviations are used to associate primary accession
 numbers given in the feature table with their source databases:
 Em: EMBL; Sw: SWISSPROT; Tr: TREMBL; Wp: WORMPEP; Information
 on the WORMPEP database can be found at
http://www.sanger.ac.uk/Projects/C_elegans/wormpep This sequence

was generated from part of bacterial clone contigs of human
 chromosome 1, constructed by the Sanger Centre Chromosome 1 Mapping
 Group. Further information can be found at
<http://www.sanger.ac.uk/HGP/Chr1>
 Rpl1-496D1 is from the library RPl1-11.2 constructed by the group
 of Pletzer de Jong. For further details see
<http://www.chori.org/bacpac/home.htm>
 VECTOR: pBAC3.6
 ----- Genome Center
 Center: Wellcome Trust Sanger Institute
 Center code: SC
 Web site: <http://www.sanger.ac.uk>
 Contact: vegas@sanger.ac.uk

 This sequence was finished as follows unless otherwise noted: all
 regions were either double-stranded or sequenced with an alternate
 chemistry or covered by high quality data (i.e., phred quality >=
 30); an attempt was made to resolve all sequencing problems, such
 as compressions and repeats; all regions were covered by at least
 one subclone; and the assembly was confirmed by restriction digest,
 except on the rare occasion of the clone being a YAC.
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KEYWORDS HTG; HTGS_PHASE1; HTGS_DRAFT; HTGS_FULLTOP.
 SOURCE Homo sapiens (human)
 ORGANISM Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini;
 Homidae; Homo.
 REFERENCE 1 (bases 1 to 195616)
 AUTHORS Birren, B., Linton, L., Nusbaum, C. and Lander, E.
 TITLE Homo sapiens chromosome 11, clone RP11-804A23
 JOURNAL Unpublished
 REFERENCE 2 (bases 1 to 195616)
 AUTHORS Birren, B., Linton, L., Nusbaum, C., Lander, E., Allen, N., Anderson, S.,
 Barina, N., Bastien, V., Boguslavsky, L., Bouckhalter, B., Brown, A.,
 Camarata, J., Campopiano, A., Colangelo, Y., Colangelo, M., Collins, S.,
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 Zembek, L., Zimmer, A. and Zody, M.
 Direct Submission
 Submitted (17-FEB-2001) Whitehead Institute/MIT Center for Genome
 Research, 320 Charles Street, Cambridge, MA 0241, USA
 On Jul 7, 2001 this sequence version replaced gi:13357354.
 All repeats were identified using RepeatMasker:
 Smit, A.F.A. & Green, P. (1996-1997)
 http://ftp.genome.washington.edu/RM/RepeatMasker.html
 ----- Genome Center
 Center: Whitehead Institute/ MIT Center for Genome Research
 Center code: WIBR
 Web site: http://www-seq.wi.mit.edu
 Contact: sequence_submissions@genome.wi.mit.edu
 ----- Project Information
 Center project name: 804_A_23
 Center clone name: L12622
 ----- Summary Statistics
 Sequencing vector: Plasmid; n/a; 100% of reads
 Chemistry: Dye-terminator Big Dye; 100% of reads
 Assembly program: Phrap; version 0.960731
 Consensus quality: 19138 bases at least Q40
 Consensus quality: 19321 bases at least Q30
 Consensus quality: 194120 bases at least Q20
 Insert size: 194716; sum-of-contigs
 Quality coverage: 9.4 in Q20 bases; sum-of-contigs

 * NOTE: This is a 'working draft' sequence. It currently
 * consists of 10 contigs. The true order of the pieces
 * is not known and their order in this sequence record is
 * arbitrary. Gaps between the contigs are represented as
 * runs of N, but the exact sizes of the gaps are unknown.
 * This record will be updated with the finished sequence
 * as soon as it is available and the accession number will
 * be preserved.
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 * 9270 9369: gap of 100 bp
 * 9370 10358: contig of 989 bp in length
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 Best Local Similarity 74.0%; Pred.No.3.5e-43;
 Matches 216; Conservative 1; Mismatches 68; Indels 7; Gaps 1;
 Oy 81 ATGCGTGAATCCAGACACTTCGGAGAGCCAAAGTGGCGCGATCACTGAGTCAAGAGA 140
 DB 91667 ATGCGTGAATCCAGACACTTCGGAGAGCCGAGCGGATCACTTGAAGTCAAGAGT 91608
 Oy 141 TCGAGACCATCTCGCCCAACATGATGAACCCCGCTTTTACTAATAAATAACAAAAATGAC 200

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Oy	201	TGGGCATGTGGGCACACACTGTATGCCAGTTACTTCAGAGA-----GCCGAGATTGC	253	
Db	91547	CAGCCGCGGGGCGCAGTTGCTCTGTAACTCCAGCTACTCCGAGACCTGAGCAGAGAAATTG	91488	
Oy	254	AGTAGCTGAAGATCGCAGAGTGAGCCGAATTCACAGATCACAGTAGCAGAGTGAGAC	313	
Db	91487	CTTTAGCCCCAGAGCGACAGAGTTGTGCACTTAAGCTTAATCTGTCTGTGGCAAGAGTGAGAC	91428	
Oy	314	KCCGCTCTCAAAAAACAACACAAAAAACAAAAAACCATTAAGACATTGTCCAT	365	
Db	91427	TCCATCTCAAGAAAAAAAAAAAAAAAAAGAAAAAGAAAAACCTGTAGTCTAT	91376	
RESULT 10 LOCUS AC090345/c				
DEFINITION	AC090345	197156 bp	DNA	linear HTG 11-JUL-2001
LOCUS	Homo sapiens chromosome 11 clone RP11-804B24 map 11,			WORKING DRAFT
SEQUENCE	SEQUENCE, 10 unordered pieces.			
VERSION	AC090345.3	GI:1467011		
KEYWORDS	HTG; HTGS_PHASE1; HTGS_DRAFT; HTGS_FULLTOP.			
SOURCE	Homo sapiens			
ORGANISM	Homo sapiens (human)			
REFERENCE	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Euarcharia; Euarchontoglires; Primates; Catarrhini; Hominae; Homo. 1 (bases 1 to 197156) Bitren,B., Linton,L., Nusbaum,C., Lander,E., Allen,N., Anderson,S., Batra,N., Bastien,V., Bonuskavsky,I., Bouckaglier,B., Brown,A., Homo sapiens chromosome 11, clone RP11-804B24 Unpublished 2 (bases 1 to 197156) Bitren,B., Linton,L., Nusbaum,C., Lander,E., Allen,N., Anderson,S., Batra,N., Bastien,V., Bonuskavsky,I., Bouckaglier,B., Brown,A., Camaretta,J., Campiano,R., Choepel,Y., Colangelo,M., Collins,S., Collamore,A., Cooke,P., Dearellano,K., Dewar,K., Diaz,J.S., Dodgson,S., Fero,S., Ferreira,P., Fitzhugh,W., Gage,D., Galagan,J., Gardyna,S., Ginde,S., Goylete,M., Graham,L., Grand-Pierre,N., Hagos,B., Heaford,A., Horton,J., Hulme,W., Iliev,I., Johnson,R., Jones,C., Karatas,A., Larocque,K., Lamazares,R., Landers,T., Lehochky,J., Levine,R., Liu,G., Maclean,C., Macdonald,P., Marquis,N., Matthews,C., McCarthy,M., McEwan,P., McKernan,K., McNeters,R., Meldrum,J., Meneses,L., Mihova,T., Mlenga,V., Murphy,T., Naylor,J., Nguyen,C., Norbu,C., Norman,C.H., O'Connor,T., O'Donnell,P., O'Neill,D., Oliver,J., Peterson,K., Phunkhang,P., Pierre,N., Pollara,V., Raymond,C., Retia,R., Rhuback,M., Riley,R., Rise,C., Rogov,P., Roman,J., Rosetti,M., Roy,A., Santos R., Schauer,S., Schnback,R., Seaman,S., Severy,P., Sougnuez,C., Spencer,B., Strange-Thomann,N., Stojanovic,C.N., Strauss,N., Subramanian,A., Talamas,J., Testaye,S., Theodore,J., Travers,M., Travis,N., Trigliolo,J., Vassiliev,H., Viel,R., Vo,A., Wilson,B., Wu,X., Wyman,D., Ye,W.J., Young,G., Zainoun,J., Zembek,L., Zimmer,A. and Zody,M.			
TITLE	Direct Submission			
JOURNAL	Submitted (17-FEB-2001) Whitehead Institute/MIT Center for Genome Research, 320 Charles Street, Cambridge, MA 02141, USA On Jul 11, 2001 this sequence version replaced g1:13357355. All repeats were identified using RepeatMasker: Smit, A.F.A. & Green, P. (1996-1997) http://ftp.genome.washington.edu/RW/RepeatMasker.html			
COMMENT	----- Genome Center Center: Whitehead Institute/ MIT Center for Genome Research Center code: WIRB Web site: http://www-seq.wi.mit.edu Contact: sequence_submissions@genome.wi.mit.edu ----- Project Information Center project name: L12623 Center clone name: 804 B 24 ----- Summary Statistics Sequencing vector: Plasmid; n/a, 100% of reads Chemistry: Dye-terminator Big Dye, 100% of reads			

Assembly program: Phrap, version 0.960731	
Consensus quality: 193219 bases at least Q40	
Consensus quality: 194831 bases at least Q30	
Consensus quality: 195508 bases at least Q20	
Insert size: 196256; sum-of-contigs	
Quality coverage: 10.0 in Q20 bases; sum-of-contigs	

* NOTE: This is a 'working draft' sequence. It currently	
* consists of 10 contigs. The true order of the pieces	
* is not known and their order in this sequence record is	
* arbitrary. Gaps between the contigs are represented as	
* runs of N, but the exact sizes of the gaps are unknown.	
* This record will be updated with the finished sequence	
* as soon as it is available and the accession number will	
* be preserved.	

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* 14985 15084: gap of 100 bp	
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* 21801 21900: gap of 100 bp	
* 21901 61396: contig of 39496 bp in length	
* 61397 61496: gap of 100 bp	
* 61497 83042: contig of 21546 bp in length	
* 83043 83143: gap of 100 bp	
* 83143 97779: contig of 14637 bp in length	
* 97780 97879: gap of 100 bp	
* 97880 124355: contig of 26476 bp in length	
* 124356 124455: gap of 100 bp	
* 124456 150156: contig of 25701 bp in length	
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Matches 216; Conservative 1; Mismatches 68; Indels 7; Gaps 1;

81 ATGCTGTAAATCCCGACGACTTGGGAGGCCAAGTGGGGGATCACTGAGGTCAAGAGA 140
DB ATGCTGTAAATCCCGACGACTTGGGAGGCCAAGTGGGGGATCACTGAGGTCAAGAGT 88321
QY 141 TCGAGACCATCTGGCCCAACATGATGTGAAACCCGCTTCTTAAATAACAAAAATTAGC 200
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QY 201 TGGGATGTGGGACACACCTGTAGTCCGACTACTCAGGA-----GCCGAGATTGC 253
DB CAGGCTGTGGGACATGCTGTATCCGACTACTCGGAGACTGAGGCAAGAAATTG 88201
QY 254 AGTGAGCTGAGATCGCAGATGAGCCGAATCAAGATCAAGACTGAGCAGAGTAGAC 313
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RESULT 11
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ACCESSION AP000777
VERSION AP000777.4 GI:28189489
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SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini;
Homnidae; Homo.

REFERENCE
AUTHORS Hattori, M., Ishii, K., Toyoda, A., Taylor, T. D., Hong-Seog, P.,
Fujiyama, A., Yada, T., Totoki, Y., Watanabe, H. and Sakaki, Y.
TITLE Homo sapiens genomic DNA
JOURNAL Published Only in Database (1999)
REFERENCE 2 (bases 1 to 197856)
AUTHORS Hattori, M., Ishii, K., Toyoda, A., Taylor, T. D., Hong-Seog, P.,
Fujiyama, A., Yada, T., Totoki, Y., Watanabe, H. and Sakaki, Y.
TITLE Direct Submission
JOURNAL Submitted (25-NOV-1999) Masahira Hattori, The Institute of Physical
and Chemical Research (RIKEN), Genomic Sciences Center (GSC),
1-7-22 Saitoh-cho, Tsukuba, Ibaraki, Japan, 305-3858, Japan
Tel: 81-45-503-9111, Fax: 81-45-503-9170
On Jan 31, 2003 this sequence version replaced gi:23821512.
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Best Local Similarity 74.0%; Pred. No. 3.5e-43;
Matches 216; Conservative 1; Mismatches 68; Indels 7; Gaps 1;

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QY 141 TCGAGACCATCTGGCCCAACATGATGTGAAACCCGCTTCTTAAATAACAAAAATTAGC 200
DB TCGAGACCATCTGGCCCAACATGATGAAACCCGCTTCTTAAATAACAAAAATTAGC 52748
QY 201 TGGGATGTGGGACACACCTGTAGTCCGACTACTCAGGA-----GCCGAGATTGC 253
DB CAGGCTGTGGGACATGCTGTATCCGACTACTCGGAGACTGAGGCAAGAAATTG 52808
QY 254 AGTGAGCTGAGATCGCAGATGAGCCGAATCAAGATCAAGACTGAGCAGAGTAGAC 313
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VERSION AC087662.1 GI:12229437
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SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini;
Homnidae; Homo.

REFERENCE
AUTHORS Birren, B., Linton, L., Nusbaum, C., Lander, E., Allen, N., Anderson, S.,
Barnett, N., Bastien, V., Boguslavsky, L., Bouckgeater, B., Brown, A.,
Camarata, J., Campopiano, A., Choquet, Y., Colangelo, M., Collins, S.,
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Souganis, C., Spencer, B., Stange-Thomann, N., Stojanovic, N.,
Straus, M., Subramanian, A., Talamas, J., Tesfaye, S., Theodore, J.,
Travers, M., Travis, N., Trigglio, J., Vassiliou, H., Viel, R., Vo, A.,
Wilson, B., Wu, X., Wyman, D., Ye, W. J., Young, G., Zainoun, D.,
Zembek, L., Zimmer, A. and Zody, M.
TITLE Direct Submission
JOURNAL Submitted (15-JAN-2001) Whitehead Institute/MIT Center for Genome
Research, 320 Charles Street, Cambridge, MA 02141, USA
COMMENT
Smit, A. F. A. & Green, P. (1996-1997)

```
http://ftp.genome.washington.edu/RM/RepeatMasker.html
----- Genome Center
Center: Whitehead Institute/ MIT Center for Genome Research
Center code: WIBR
Web site: http://www-seq.wi.mit.edu
Contact: sequence.submissions@genome.wi.mit.edu
----- Project Information
Center project name: L12144
Center clone name: 1155_K23

* NOTE: This record contains 84 individual
* sequencing reads that have not been assembled into
* contigs. Runs of N are used to separate the reads
* and the order in which they appear is completely
* arbitrary. Low-pass sequence sampling is useful for
* identifying clones that may be gene-rich and allows
* overlap relationships among clones to be deduced.
* However, it should not be assumed that this clone
* will be sequenced to completion. In the event that
* the record is updated, the accession number will
* be preserved.
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* 710
* 809: gap of 100 bp
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* 1546: contig of 737 bp in length
* 1547
* 1646: gap of 100 bp
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* 2377: contig of 731 bp in length
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* 2477: gap of 100 bp
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* 3197: contig of 720 bp in length
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* 3297: gap of 100 bp
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* 4024: contig of 727 bp in length
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* 4124: gap of 100 bp
* 4125
* 4842: contig of 718 bp in length
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* 4943
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* 7418
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* 14812
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* 16323: contig of 698 bp in length
* 16324
* 16423: gap of 100 bp
* 16424
* 17156: contig of 733 bp in length
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* 20548
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* 22085: contig of 718 bp in length
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* 22883: contig of 698 bp in length
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* 47622: contig of 716 bp in length
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* 51712 51811: gap of 100 bp
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 * 54151 54250: gap of 100 bp
 * 54251 54967: contig of 717 bp in length
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Query Match 39.7%; Score 158.6; DB 12; Length 66990;
 Best Local Similarity 77.7%; Pred. No. 2.5e-43;

Matches 206; Conservative 1; Mismatches 50; Indels 8; Gaps 1;

84 CCTGTATCCAGCATTGGGAGGCCAAGTGGCGGATCACTGAGGTCAAGATCG 143
 32552 COTGTATCCAGCATTGGGAGGCCAAGTGGCGGATCACTGAGGTCAAGATCG 32493

144 AGACCATCTGGCCCACTGTGAAACCCGCTTTACTAAATAACAAAATAGCTGG 203
 32492 ATACCAAGCTGGCCCACTGTGAAACCCGCTTTACTAAATAACAAAATAGCTGG 32433

204 GCATGTGGGACACACCTGTAGTCCGACTACTCAGAGCCGGAATGGCATGCTGA 263
 32432 GCGTGTGTGTGACCGCTATATATCCGACTACTCAGAGCCGGAATGGCATGCTGA 32373

264 GATCGCAGAGTGGCCGAATATCATCATGATCAGATGAGAGTGAAGACCCGCTCA 323
 32372 GATTG-----TGCATTTGCACCGCTAGCCCTGGGCAACAGAGTGAATCTATCTCA 32321

324 AAACACACACAAAAACAAAAAAC 348
 32320 AAAAAAAAAAAAAAAAAAATATATAC 32296

RESULT 13
 DD164421 96593 bp DNA linear PAT 23-NOV-2005

LOCUS DD164421 96593 bp DNA linear PAT 23-NOV-2005
 DEFINITION NOVEL COMPOSITIONS AND METHODS FOR CANCER.
 ACCESSION DD164421
 VERSION DD164421.1 GI:83948746
 KEYWORDS JP 2005510225-A/16.
 SOURCE unidentified
 ORGANISM unidentified
 unclassified sequences.
 1 (bases 1 to 96593)

REFERENCE 1 (bases 1 to 96593)
 MAURICE,D.W.
 NOVEL COMPOSITIONS AND METHODS FOR CANCER
 TITLE Patent: JP 2005510225-A 16 21-APR-2005;
 JOURNAL SAGES DISCOVERY
 OS Homo Sapien
 PN JP 2005510225-A/16
 PD 21-APR-2005
 PF 02-DEC-2002 JP 2003546739
 PR 30-NOV-2001 US 09/997722
 PI david w maurice
 CC

COMMENT
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 /organism="unidentified"
 /mol_type="unassigned DNA"
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ORIGIN
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 Best Local Similarity 75.0%; Pred. No. 3.4e-43;
 Matches 216; Conservative 1; Mismatches 57; Indels 14; Gaps 1;

81 ATGCTGTATCCAGCATTGGGAGGCCAAGTGGCGGATCACTGAGGTCAAGGA 140

18217 ACGCTGTATCCAGCATTGGGAGGCCAAGTGGCGGATCACTGAGGTCAAGGA 18276

141 TCGAGACCATCTGGCCCACTGTGAAACCCGCTTTACTAAATAACAAAATAGC 200
 18277 TCAAGATCAGCTGTGGCCCACTGTGAAACCCGCTTTACTAAATAACAAAATAGC 18336

201 TGGGCATGTGGGACACACCTGTAGTCCAGCTTACTCAGAGCCGGAATGGCATGAGC 260
 18337 TGGGCATGTGGGACACACCTGTAGTCCAGCTTACTCAGAGCCGGAATGGCATGAGC 18396

261 TGAATCGCAG-----AGTAGCCGCAATTCACAGATCACAAGTGAAGCA 306
 18397 CTTAATCCAGGCGGCGGAGGTTCAGATGAGCCAGATTCGACAGCTTGGGTGACAGC 18456

307 GTGAGACCGCTCTCAAAAACACACAAAAACAAAACCAATAG 354
 18457 GTGAGACTTATCTCAAAAACACACAAAAACAAAACCAATAG 18504

RESULT 15
 AC010677/c 112659 bp DNA linear PRI 03-JAN-2002

LOCUS AC010677/c 112659 bp DNA linear PRI 03-JAN-2002
 DEFINITION Homo sapiens chromosome 7 clone CTD-230414, complete sequence.

Query Match 39.6%; Score 158.4; DB 2; Length 96593;
 Best Local Similarity 75.0%; Pred. No. 3.4e-43;
 Matches 216; Conservative 1; Mismatches 57; Indels 14; Gaps 1;

81 ATGCTGTATCCAGCATTGGGAGGCCAAGTGGCGGATCACTGAGGTCAAGGA 140

18217 ACGCTGTATCCAGCATTGGGAGGCCAAGTGGCGGATCACTGAGGTCAAGGA 18276

141 TCGAGACCATCTGGCCCACTGTGAAACCCGCTTTACTAAATAACAAAATAGC 200
 18277 TCAAGATCAGCTGTGGCCCACTGTGAAACCCGCTTTACTAAATAACAAAATAGC 18336

201 TGGGCATGTGGGACACACCTGTAGTCCAGCTTACTCAGAGCCGGAATGGCATGAGC 260
 18337 TGGGCATGTGGGACACACCTGTAGTCCAGCTTACTCAGAGCCGGAATGGCATGAGC 18396

261 TGAATCGCAG-----AGTAGCCGCAATTCACAGATCACAAGTGAAGCA 306
 18397 CTTAATCCAGGCGGCGGAGGTTCAGATGAGCCAGATTCGACAGCTTGGGTGACAGC 18456

307 GTGAGACCGCTCTCAAAAACACACAAAAACAAAACCAATAG 354
 18457 GTGAGACTTATCTCAAAAACACACAAAAACAAAACCAATAG 18504

RESULT 14
 AK695641 96593 bp DNA linear PAT 31-MAR-2003

LOCUS AK695641 96593 bp DNA linear PAT 31-MAR-2003
 DEFINITION Sequence 1268 from Patent WO03008583.
 ACCESSION AK695641
 VERSION AK695641.1 GI:29418793
 KEYWORDS Homo sapiens (human)
 SOURCE Homo sapiens
 ORGANISM Homo sapiens
 unclassified sequences.
 1 (bases 1 to 96593)

REFERENCE 1
 MORRIS,D.W. and Engelhard,E.K.
 Novel compositions and methods for cancer
 TITLE Patent: WO 03008583-A 1268 30-JAN-2003;
 JOURNAL Sages Discovery (US)
 OS Homo Sapien
 PN WO 03008583-A 1268 30-JAN-2003;
 PD 30-JAN-2003
 PF 03-JAN-2002
 PR 03-JAN-2002
 PI 03-JAN-2002
 CC

COMMENT
 FH Key Location/Qualifiers.
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ORIGIN
 Query Match 39.6%; Score 158.4; DB 2; Length 96593;
 Best Local Similarity 75.0%; Pred. No. 3.4e-43;
 Matches 216; Conservative 1; Mismatches 57; Indels 14; Gaps 1;

81 ATGCTGTATCCAGCATTGGGAGGCCAAGTGGCGGATCACTGAGGTCAAGGA 140

ACCESSION	AC010677
VERSION	AC010677.4
KEYWORDS	GI:11465112
SOURCE	HTG.
ORGANISM	Homo sapiens (human)
REFERENCE	Homo sapiens Embryonic; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini; Hominidae; Homo
AUTHORS	1 (bases 1 to 112659) Waterston,R.H.
TITLE	The sequence of Homo sapiens c1one
JOURNAL	Unpublished
AUTHORS	2 (bases 1 to 112659) Waterston,R.H.
TITLE	Direct Submission
JOURNAL	Submitted (17-SEP-1999) Genome Sequencing Center, Washington University School of Medicine, 4444 Forest Park Parkway, St. Louis, MO 63108, USA
REFERENCE	3 (bases 1 to 112659) Waterston,R.H.
AUTHORS	Direct Submission
TITLE	Submitted (30-NOV-2000) Genome Sequencing Center, Washington University School of Medicine, 4444 Forest Park Parkway, St. Louis, MO 63108, USA
JOURNAL	4 (bases 1 to 112659) Waterston,R.H.
REFERENCE	Direct Submission
AUTHORS	Submitted (09-MAY-2001) Department of Genetics, Washington University, 4444 Forest Park Avenue, St. Louis, Missouri 63108, USA
TITLE	5 (bases 1 to 112659) Waterston,R.H.
JOURNAL	Direct Submission Submitted (03-JUN-2002) Genome Sequencing Center, Washington University School of Medicine, 4444 Forest Park Parkway, St. Louis, MO 63108, USA
COMMENT	On Nov 30, 2000 this sequence version replaced gi:7630788.
FEATURES	----- Genome Center ----- Center: Washington University Genome Sequencing Center Center code: MUGSC Web site:http://genome.wustl.edu/gsc/index.shtml Contact: submissions@watsen.wustl.edu ----- Project Information ----- Center project name: H_MS2304L04 ----- Location/Qualifiers 1. 112659 /organism="Homo sapiens" /mol_type="genomic DNA" /db_xref="taxon:9606" /chromosome="7" /clone="CTD-2304L4"
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Best Local Similarity	73.9%; Pred. No. 3,7e-43;
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QY	136 AGAGATCGAGACCATCTCTGGCGCAACATGATGAAACCCCGCTTTACTAATAAATACAAAA 195
Db	16899 GGAATTGAGATCAAGTCTGGCGCAACATGATGAAACCCCGCTTTACTAATAAATACAAAA 168400
QY	196 ATAGCTGGGCGATGGTGGCAACACCTGTAGTCCAGCTACTCAAGAGCGGAGATTGGCAG 255
Db	16839 CCAGCTGGGTGTGGTGAACGCGTTGTAATCCAGCTACTTGGAGGCGGAGATTGGCAG 167800
QY	256 TGAAGTCGAGATCGGAGAGTGAAGCCGAATACAGATACAGATGAGGACAGAGTGAACKC 315
Db	16779 TGAAGTCGAGATCGGAGATCACTACCTCAGCGCTGGGTGAACAGAGGAGATCTGTCTCCA 167200

Oy		316	CGCTCAAAAACAAACAACAAAAA	347
Dd		16719	GGAAAAAAAAAAAAAAAAAAAAAA	16688
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VERSION		AC026407.4		GI:15375158
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SOURCE		Homo sapiens (human)		
ORGANISM		Homo sapiens		Eukaryote; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini; Homidae; Homo.
REFERENCE		1 (bases 1 to 155344)		DOE Joint Genome Institute and Stanford Human Genome Center.
AUTHORS		Direct Submission		
TITLE		Unpublished		
JOURNAL		2 (bases 1 to 155344)		
REFERENCE		DOE Joint Genome Institute.		
AUTHORS		Direct Submission		
TITLE		Submitted (22-MAR-2000) Production Sequencing Facility, DOE Joint		
JOURNAL		Genome Institute, 2800 Mitchell Drive, Walnut Creek, CA 94598, USA		
REFERENCE		3 (bases 1 to 155344)		
AUTHORS		DOE Joint Genome Institute and Stanford Human Genome Center.		
TITLE		Direct Submission		
JOURNAL		Submitted (18-JUL-2001) DOE Joint Genome Institute, 2800 Mitchell		
REFERENCE		Drive, Walnut Creek, CA 94598, USA		
AUTHORS		4 (bases 1 to 155344)		
TITLE		DOE Joint Genome Institute and Stanford Human Genome Center.		
JOURNAL		Direct Submission		
REFERENCE		Submitted (30-AUG-2001) DOE Joint Genome Institute, 2800 Mitchell		
AUTHORS		Drive, Walnut Creek, CA 94598, USA		
TITLE		On Aug 30, 2001 this sequence version replaced gi:14861724.		
JOURNAL		Draft Sequence Produced by DOE Joint Genome Institute		
COMMENT		www.jgi.doe.gov		
		Finishing Completed at Stanford Human Genome Center		
		www.sngc.stanford.edu		
		Quality: Phrap Quality >=40 99.8% of Sequence;		
		Estimated Total Number of Errors is 0.2.		
		SMS Content:		
		WI-11894 G21380		
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Best Local Similarity		78.5%;	Pred. No. 5.1e-43;	
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Dd		95307	ATGCTTTAATCCAGACTTTTGGAAGGCTGAGAGCAGGGGAAATTACTAGGCTGGAAAT	95366
Oy		141	TCGAGACCATCTCGGCCAACATGCTGAACCCCGTTTACTTAAATAATCAAAAAATAGC	200
Dd		95367	TCAAGACCAACCTGGCCCAATGATAAAACCCGTCTTACTTAAAAAATCAAAAAATCACG	95428
Oy		201	TGGGCAATGGGAGCACAACCTGATAGTCCAGAGTCACTCAGAGCGGAGATTGCAATGAGC	260
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QY 261 TGAGATCGCAGAGTGGCCGAATATCAGATCAGAGAGGAGAGTGAGACXCGCTCT 320
 DB 95487 TGAAGTTG-----TGCACCTGCACTTCAGAGCTGGGTGACAAAGCAAACTGTCT 95538
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 DB 95539 CAAAAAAAAAAAAAAAAAGCA 95559
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 LOCUS Homo sapiens chromosome 16 clone RP11-46107, complete sequence.
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 AC040168
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 SOURCE Homo sapiens (human)
 ORGANISM Homo sapiens
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 Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini;
 Homnidae; Homo.
 REFERENCE 1 (bases 1 to 170154)
 AUTHORS Doe Joint Genome Institute, Stanford Human Genome Center and Los
 Alamos National Laboratory.
 TITLE Direct Submission
 JOURNAL Unpublished
 REFERENCE 2 (bases 1 to 170154)
 AUTHORS Doe Joint Genome Institute.
 TITLE Direct Submission
 JOURNAL Submitted (11-APR-2000) Production Sequencing Facility, DOE Joint
 Genome Institute, 2800 Mitchell Drive, Walnut Creek, CA 94598, USA
 REFERENCE 3 (bases 1 to 170154)
 AUTHORS Doe Joint Genome Institute.
 TITLE Direct Submission
 JOURNAL Submitted (03-MAY-2002) Production Sequencing Facility, DOE Joint
 Genome Institute, 2800 Mitchell Drive, Walnut Creek, CA 94598, USA
 REFERENCE 4 (bases 1 to 170154)
 AUTHORS Doe Joint Genome Institute.
 TITLE Direct Submission
 JOURNAL Submitted (22-JAN-2003) Production Sequencing Facility, DOE Joint
 Genome Institute, 2800 Mitchell Drive, Walnut Creek, CA 94598, USA
 REFERENCE 5 (bases 1 to 170154)
 AUTHORS Doe Joint Genome Institute, Stanford Human Genome Center and Los
 Alamos National Laboratory.
 TITLE Direct Submission
 JOURNAL Submitted (29-MAR-2003) DOE Joint Genome Institute, 2800 Mitchell
 Drive, Walnut Creek, CA 94598, USA
 COMMENT On Mar 29, 2003 this sequence version replaced gi:27819471.
 Draft Sequence Produced by DOE Joint Genome Institute
 www.jgi.doe.gov
 Finishing Completed at Stanford Human Genome Center and Los Alamos
 National Laboratory
 www.sbgc.stanford.edu
 Quality: Phrap Quality >=40 100% of Sequence;
 Estimated Total Number of Errors is 0.
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QY 141 TCGAGACCATCTGSCCAACATGTGTAACCCCGTCTTAACTAATAAATACAAATATAGC 200
 DB 134379 TCGAGACCATCTGSCCAACATGTGTAACCCCGTCTTAACTAATAAATACAAATATAGC 134438
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 QY 321 CAAAACACACACAAACAAACAAACACATPA 353
 DB 134559 CAAAAAAAAAAAAAAAAACCAAAAAAAAAAGGA 134591
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 DEFINITION AC018605
 AC018605
 VERSION AC018605.4 GI:10047801
 KEYWORDS HTG; HTGS PHASE1; HTGS_DRAFT.
 SOURCE Homo sapiens (human)
 ORGANISM Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini;
 Homnidae; Homo.
 REFERENCE 1 (bases 1 to 186747)
 AUTHORS Birren, B., Linton, L., Nusbaum, C. and Lander, E.
 Homo sapiens chromosome 16, clone RP11-764C24
 Unpublished
 REFERENCE 2 (bases 1 to 186747)
 AUTHORS Birren, B., Linton, L., Nusbaum, C., Lander, E., Abrahams, H., Allen, N.,
 Anderson, S., Baldwin, J., Barna, N., Beckler, R., Beda, F.,
 Boguslavskiy, L., Boukhgalter, B., Brown, A., Burkett, G., Castele, A.,
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 Santos, R., Severy, P., Spencer, B., Stange-Thomann, N., Stojanovic, N.,
 Subramanian, A., Talamas, J., Testaye, S., Theodore, J., Tirrell, A.,
 Vassiliev, H., Viel, R., Vo, A., Wu, X., Wyman, D., Ye, W. J., Zimmer, A.
 and Zody, M.
 Direct Submission
 JOURNAL Submitted (14-DEC-1999) Whitehead Institute/MIT Center for Genome
 Research, 320 Charles Street, Cambridge, MA 02141, USA
 REFERENCE 3 (bases 1 to 186747)
 AUTHORS Birren, B., Linton, L., Nusbaum, C., Lander, E., Abrahams, H., Allen, N.,
 Anderson, S., Barna, N., Bastien, V., Beda, F., Boguslavskiy, L.,
 Boukhgalter, B., Brown, A., Burkett, G., Campion, A., Castele, A.,
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 MacDonald, P., Marquis, N., McCarty, M., McEwan, P., McKernan, K.,
 McHeeters, R., Meldrim, J., Menus, L., Mihova, T., Mlenga, V.,
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 O'Donnell, P., O'Neill, D., Olivari, T. M., Oliver, J., Peterson, K.,
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 Rogov, P., Rothman, D., Roy, A., Santos, R., Schauer, S., Severy, P.,
 Stojanovic, N., Spencer, B., Stange-Thomann, N., Stojanovic, N.,
 Straube, N., Subramanian, A., Talamas, J., Testaye, S., Theodore, J.,

TITLE
JOURNAL
COMMENT

Tirelli, A., Travers, M., Trigilio, J., Vaassiliev, H., Viel, R., Vo, A., Wilson, B., Wu, X., Wyman, D., Ye, W. J., Young, G., Zainoun, J., Zimmer, A. and Zody, M.
Submitted (24-AUG-2002) Whitehead Institute/MIT Center for Genome Research, 320 Charles Street, Cambridge, MA 02141, USA
On Sep 10, 2000 this sequence version replaced gi:7407949.
All repeats were identified using RepeatMasker:
Smit, A.F.A. & Green, P. (1996-1997)
<http://ftp.genome.washington.edu/RM/RepeatMasker.html>

Center: Whitehead Institute/ MIT Center for Genome Research

Center code: MIBR

Web site: <http://www-seq.wi.mit.edu>

Contact: sequence.submissions@genome.wi.mit.edu

Project Information

Center project name: L4986

Center clone name: 764_C_24

Summary Statistics

Sequencing vector: M13; M7815; 100% of reads
Chemistry: Dye-terminator Big Dye; 100% of reads
Assembly program: Phrap; version 0.960731

Consensus quality: 174578 bases at least Q40

Consensus quality: 179739 bases at least Q30

Consensus quality: 182173 bases at least Q20

Insert size: 188000; agarose-fp

Insert size: 185347; sum-of-contigs

Quality coverage: 4.6 in Q20 bases; agarose-fp

Quality coverage: 4.7 in Q20 bases; sum-of-contigs

* NOTE: This is a 'working draft' sequence. It currently
* consists of 15 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence
* as soon as it is available and the accession number will
* be preserved.

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* 26749 26848: gap of 100 bp
* 26849 28315: contig of 1467 bp in length
* 28316 28415: gap of 100 bp
* 28416 30899: contig of 2484 bp in length
* 30900 30999: gap of 100 bp
* 31000 34327: contig of 3328 bp in length
* 34328 34427: gap of 100 bp
* 34428 39283: contig of 4856 bp in length
* 39284 39383: gap of 100 bp
* 39384 44613: contig of 5210 bp in length
* 44614 44713: gap of 100 bp
* 44714 78567: contig of 33854 bp in length
* 78568 78667: gap of 100 bp
* 78668 89039: contig of 10372 bp in length
* 89040 89139: gap of 100 bp
* 89140 97713: contig of 8574 bp in length
* 97714 97813: gap of 100 bp
* 97814 110219: contig of 12406 bp in length
* 110220 110319: gap of 100 bp
* 110320 123656: contig of 13337 bp in length
* 123657 123756: gap of 100 bp
* 123757 138462: contig of 14706 bp in length
* 138463 138562: gap of 100 bp
* 138563 153566: contig of 15004 bp in length
* 153567 153666: gap of 100 bp
* 153667 180503: contig of 26837 bp in length
* 180504 180604: gap of 100 bp
* 180604 186747: contig of 6144 bp in length.

FEATURES
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/db_xref="taxon:9606"
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Best Local Similarity 75.8%; Pred. No. 7.6e-43;
Matches 207; Conservative 1; Mismatches 63; Indels 2; Gaps 1;
81 ATGCGTGAATCCAGCACTTCGGAGAGCGCAAGTGGCGGATCACTGAGTCAAGAGA 140
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Db      69948  ACGCTGTATATCCAGCACTTTGGAGGCGCAAGTGGGAGATCNC--GAGGTCAAGGA 70005
Qy      141  TCGAGACCATCTGGCCCAACATGTGAAACCCCGTCTTACTTAAATAACAAAAATAGC 200
Db      70006  TCGAGACCATCTGGCCCAACATGTGAAACCCCGTCTTACTTAAATAACAAAAATAGC 70065
Qy      201  TGGGATGTGGGACACACCTGTAGTCCAGCTACTCAGAGCCGAGATTGCACTGAGC 260
Db      70066  CGGGCGTGTGGCTGGGACCTGTATATCCAGCTACTTGGAGCTGAGGCAAGATATCG 70125
Qy      261  TGAATTCGACAGTGAAGCCGAATACAGATCAGATGAGTGAAGTGAACKCCGTCT 320
Db      70126  CTTGAACCTGGAGGCGGAGGTGCACTCCAGCTGTGACAGAGAGACTCCGCT 70185
Qy      321  CAAAAACAACAACAAAAACAAACCATTA 353
Db      70186  CAAAAACAAAAACAAAAACCAAAAAAGAA 70218

RESULT 19
LOCUS      AC007495      198575 bp      DNA      linear      PRI 22-MAR-2003
DEFINITION Homo sapiens chromosome 16 clone RP11-355E10, complete sequence.
ACCESSION  AC007495
VERSION     AC007495.9  GI:29150335
KEYWORDS    HTG.

SOURCE      Homo sapiens (human)
ORGANISM    Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini;
Hominae; Homo.

REFERENCE   1 (bases 1 to 198575)
AUTHORS    DOE Joint Genome Institute, Stanford Human Genome Center and Los
            Alamos National Laboratory.
TITLE       Unpublished
JOURNAL     2 (bases 1 to 198575)
AUTHORS     Bruce, D., Mundt, M., Doggett, N., Munk, C., Saunders, E., Robinson, D.,
            Jones, M., Buckingham, J., Chasteen, L., Thompson, S., Goodwin, L.,
            Bryant, J., Tesmer, J., Meincke, L., Longmire, J., White, S., Tatum, O.,
            Campbell, C., Fawcett, J., Maltbie, M., Buesod, M., Sutherland, R.,
            McMurtry, K., Han, C. and Deaven, L.
TITLE       Direct Submission
JOURNAL     Submitted (06-MAY-1999) Center for Human Genome Studies, DOE Joint
            Genome Institute, Los Alamos National Laboratory, MS M888, Los
            Alamos, NM 87545, USA
            3 (bases 1 to 198575)
            DOE Joint Genome Institute.
REFERENCE   4 (bases 1 to 198575)
AUTHORS    DOE Joint Genome Institute.
TITLE       Direct Submission
JOURNAL     Submitted (06-FEB-2002) Production Sequencing Facility, DOE Joint
            Genome Institute, 2800 Mitchell Drive, Walnut Creek, CA 94598, USA
            5 (bases 1 to 198575)
            DOE Joint Genome Institute, Stanford Human Genome Center and Los
            Alamos National Laboratory.
TITLE       Direct Submission
JOURNAL     Submitted (22-MAR-2003) DOE Joint Genome Institute, 2800 Mitchell
            Drive, Walnut Creek, CA 94598, USA
            On Mar 22, 2003 this sequence version replaced gi:27363205.
COMMENT     Draft Sequence Produced by DOE Joint Genome Institute
            www.jgi.doe.gov
            Finishing Completed at Stanford Human Genome Center and Los Alamos
            National Laboratory
            www-shgc.stanford.edu
            Quality: Phrap Quality >=40 100% of Sequence;
            Estimated Total Number of Errors is 0.
            NOTE: BACTERIAL TRANSPOSON excised at 8530.
            Location/Qualifiers
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ORIGIN

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Best Local Similarity 75.8%; Pred. No. 7.9e-43;
Matches 207; Conservative 1; Mismatches 63; Indels 2; Gaps 1;

Qy      81  ATGCTGTATATCCAGCACTTGGAGGCGCAAGTGGGAGATCAGCTGAGTCAAGGA 140
Db      108130  ACGCTGTATATCCAGCACTTGGAGGCGCAAGTGGGAGATCAGCTGAGTCAAGGA 108187
Qy      141  TCGAGACCATCTGGCCCAACATGTGAAACCCCGTCTTACTTAAATAACAAAAATAGC 200
Db      108188  TCGAGACCATCTGGCCCAACATGTGAAACCCCGTCTTACTTAAATAACAAAAATAGC 108247
Qy      201  TGGGATGTGGGACACACCTGTAGTCCAGCTACTCAGAGCCGAGATTGCACTGAGC 260
Db      108248  CGGGCGTGTGGCTGGGACCTGTATATCCAGCTACTTGGAGCTGAGGCAAGATATCG 108307
Qy      261  TGAATTCGACAGTGAAGCCGAATACAGATCAGATGAGTGAAGTGAACKCCGTCT 320
Db      108308  CTTGAACCTGGAGGCGGAGGTGCACTCCAGCTGTGACAGAGAGACTCCGCT 108367
Qy      321  CAAAAACAACAACAAAAACAAACCATTA 353
Db      108368  CAAAAACAAAAACAAAAACCAAAAAAGAA 108400

RESULT 20
LOCUS      AL732374      224187 bp      DNA      linear      PRI 18-MAY-2005
DEFINITION Human DNA sequence from clone RP13-444X19 on chromosome X contains
            a mitochondrial ribosomal protein S18C (MRPS18C) pseudogene, the 3'
            end of the gene for a novel protein similar to PHD finger protein 2
            PHF2 and a CpG island, complete sequence.
ACCESSION  AL732374
VERSION     AL732374.14  GI:23476649
KEYWORDS    HTG; CpG island; MRPS18C; PHF2.

SOURCE      Homo sapiens (human)
ORGANISM    Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini;
Hominae; Homo.

REFERENCE   1 (bases 1 to 224187)
AUTHORS    Chapman, J.
TITLE       Direct Submission
JOURNAL     Submitted (13-MAY-2005) Wellcome Trust Sanger Institute, Hinxton,
            Cambridgeshire, CB10 1SA, UK. E-mail enquiries: vegas@sanger.ac.uk
            Clone requests: clonerequests@sanger.ac.uk
            On Oct 2 2002 this sequence version replaced gi:23393869.
            The following abbreviations are used to associate primary accession
            numbers given in the feature table with their source databases:
            Em: EMBL; Sw: SWISSPROT; Tr: TREMBL; Wp: WORMPEP; Information
            can be found at
            http://www.sanger.ac.uk/Projects/C_elegans/wormpep This sequence
            was generated from part of bacterial clone contigs of human
            chromosome X, constructed by the Sanger Centre Chromosome X Mapping
            Group. Further information can be found at
            http://www.sanger.ac.uk/HGP/Chix
            RP13-444X19 is from the library RP13-13.2 constructed by the group
            of Pieter de Jong. For further details see
            http://www.choxi.org/bacpac/home.htm
            VECTOR: pBACE3.6
            ----- Genome Center
            Center: Wellcome Trust Sanger Institute
            Center code: SC
            Web site: http://www.sanger.ac.uk
            Contact: vegas@sanger.ac.uk
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 Best Local Similarity 70.0%; Pred. No. 8.3e-43;
 Matches 229; Conservative 1; Mismatches 88; Indels 9; Gaps 1;

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 DB 190879 TCCGAGCACTTCGGAGGCGCAAGTGGCGGATCACTGAGTCAAGATCGAGCAT 190938
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 DB 190939 CATGGCCAAATGTGTGAATACCCCGTCTGTACTAAATAAATAAATAATGCGAGGATGT 190998
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 DB 190999 GGCAACACCTGTGTGTCTCCACTACTCGAGCGCGGAGATTCGACTGCTGAGTCCGA 191058
 QY 271 GAGTGGAGCGCAATCACAGATGACAGATGAGAGAGACGCCGCTCAAAAAACAC 330
 DB 191059 CCATGACACTCA-----GCCTGAGCAACAGAGTGAACATCTCTCTCAAAAAAAA 191109
 QY 331 AACAAAAACAAAAAACCATAGACA 357
 DB 191110 AAAGAAAGAAAGAAAGAAAGAAAA 191136

RESULT 21
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 LOCUS
 DEFINITION
 AC150910 169908 bp DNA linear HTG 09-OCT-2004
 Pan troglodytes chromosome 7 clone CH251-484K21, WORKING DRAFT
 SEQUENCE 8 unordered pieces.

ACCESSION
 AC150910 GI:54019620
 VERSION
 HTG; HTGS PHASE1; HTGS DRAFT; HTGS_FULUTOP.
 KEYWORDS
 Pan troglodytes (chimpanzee)
 SOURCE
 Pan troglodytes
 ORGANISM
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Butheria; Euarchontoglires; Primates; Catarrhini;
 Homiidae; Pan.
 1 (bases 1 to 169908)
 Wilson, R.K.
 The sequence of Pan troglodytes clone
 Unpublished
 2 (bases 1 to 169908)
 Wilson, R.K.
 Direct Submission
 Submitted (14-AUG-2004) Genetics, Genome Sequencing Center, 4444
 Forest Park Parkway, St. Louis, MO 63108, USA
 3 (bases 1 to 169908)
 Wilson, R.K.
 Direct Submission
 Submitted (09-OCT-2004) Genetics, Genome Sequencing Center, 4444
 Forest Park Parkway, St. Louis, MO 63108, USA
 JOURNAL
 COMMENT
 On Oct 9, 2004 this sequence version replaced gi:51241995.

Center: Washington University Genome Sequencing Center
 Center code: WUGSC
 Web site: <http://genome.wustl.edu>
 Contact: submissions@watson.wustl.edu
 Project Information
 Center project name: C_AB0484K21
 Summary Statistics
 Sequencing vector: M13, 0%
 Sequencing vector: plasmid, 100%
 Chemistry: Dye-primer ET; 0% of reads
 Chemistry: Dye-terminator Big Dye; 100% of reads
 Assembly program: Phrap; version 0.990319
 Consensus quality: 164720 bases at least Q40
 Consensus quality: 165787 bases at least Q30
 Consensus quality: 165527 bases at least Q20
 NOTE: This is a 'working draft' sequence. It currently
 * consists of 8 contigs. The true order of the pieces
 * is not known and their order in this sequence record is
 * arbitrary. Gaps between the contigs are represented as
 * runs of N, but the exact sizes of the gaps are unknown.
 * This record will be updated with the finished sequence
 * as soon as it is available and the accession number will
 * be preserved.

1 1381: contig of 1381 bp in length
 * 1382 1481: gap of unknown length
 * 1482 7431: contig of 5950 bp in length
 * 7432 7531: gap of unknown length
 * 7532 21676: contig of 14145 bp in length
 * 21677 21776: gap of unknown length
 * 21777 34793: contig of 13017 bp in length
 * 34794 34893: gap of unknown length
 * 34894 54353: contig of 19460 bp in length
 * 54354 54453: gap of unknown length
 * 54454 88284: contig of 33831 bp in length
 * 88285 88384: gap of unknown length
 * 88385 125437: contig of 37053 bp in length
 * 125438 125537: gap of unknown length
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SEQUENCING READ COVERAGE: Sequencing is completed to a minimum standard of double strand coverage with a minimum of 2 clones and 2 reads with no ambiguities or 2 chemistries with a minimum of 2 clones and 3 reads with no ambiguities. If the sequence quality for a region does not meet this standard, it will be indicated in the annotation as Low Coverage.

QUALITY OF INDIVIDUAL BASES: This sequence meets stringent quality standards - estimated error rate less than 1 per 10,000 bases. Reports of lowest quality individual bases and measures of base quality are listed below. Description of the metrics can be found at URL:

http://www.hgsc.bcm.tmc.edu:8088/quality.info/genbank.annotation.html.

FEATURES

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repeat_region
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repeat_region
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repeat_region
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Query Match 39.4%; Score 157.4; DB 5; Length 174097;
 Best Local Similarity 75.8%; Pred. No. 1e-42;
 Matches 207; Conservative 1; Mismatches 62; Indels 3; Gaps 1;

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QY 81 ATGCTGTAAATCCAGACCTTCGGAGGCGCAAGGTGGCGGATACCTGAGGTCAAGGA 140
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QY 141 TCGAGACCATCTCGGCAACATGTGTAACCCCTCTTACTAAAAATCAAAAAATAGC 200
DB 4274 TTGAGACACAGCTCGGCAACATGTGTAACCCCTCTTACTAAAAATCAAAAAATAGC 4333
QY 201 TGGGATGTGTGGCACACCTGTGTGTCCTGACTGACTGAGAGCCGAGATGTCAGTGC 260
DB 4334 TGGGATGTGTGGCACACCTGTGTGTCCTGACTGACTGAGAGCCGAGATGTCAGTGC 4393
QY 261 TGAGATCGCAGAGAGAGCCGAAATCAAGATCAAGATGTCAGTGC 317
DB 4394 CTTGAATCCGAGAGAGAGATGATGATGATGATGATGATGATGATGATGATGATG 4453
QY 318 TCTCAAAACCAACCAACCAACCAACCAACCAACCAACCAACCAACCAACCAACCA 350
DB 4454 TCTCAAAACCAACCAACCAACCAACCAACCAACCAACCAACCAACCAACCAACCA 4486

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RESULT 23
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 LOCUS Homo sapiens chromosome 19 clone CTF-33G10, complete sequence.
 AC011495
 DEFINITION AC011495.8 GI:21747443
 VERSION AC011495.8
 KEYWORDS HTG.
 SOURCE Homo sapiens (human)
 ORGANISM Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini;
 Hominidae; Homo.
 REFERENCE 1 (bases 1 to 209844)
 AUTHORS DOE Joint Genome Institute and Stanford Human Genome Center.
 TITLE Direct Submission
 JOURNAL Unpublished
 REFERENCE 2 (bases 1 to 209844)
 AUTHORS DOE Joint Genome Institute.
 TITLE Direct Submission
 JOURNAL Submitted (07-OCT-1999) Production Sequencing Facility, DOE Joint
 Genome Institute, 2800 Mitchell Drive, Walnut Creek, CA 94598, USA
 REFERENCE 3 (bases 1 to 209844)
 AUTHORS DOE Joint Genome Institute and Stanford Human Genome Center.

Direct Submission
Submitted (30-JUL-2000) Genome Sequencing Center, Washington
University School of Medicine, 4444 Forest Park Parkway, St. Louis
MO 63108, USA
On Aug 4, 2000 this sequence version replaced gi:9587428.

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Genome Center -----
Center: Washington University Genome Sequencing Center
Center code: WUGSC
Web site: http://genome.wustl.edu/gsc/index.shtml
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Project Information -----
Center project name: H_NH0507P19
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Summary Statistics -----
Sequencing vector: M13; 100%
Sequencing vector: plasmid; 0%
Chemistry: Dye-terminator Big_Dye; 0% of reads
Chemistry: Dye-terminator Big_Dye; 0% of reads
Assembly program: Phrap; version 0.990319
Consensus quality: 158826 bases at least Q40
Consensus quality: 15318 bases at least Q30
Consensus quality: 165604 bases at least Q20
Insert size: 16000; agarose-fp
Insert size: 171556; sum-of-contigs
Quality coverage: 4.60 in Q20 bases; agarose-fp
Quality coverage: 4.61 in Q20 bases; sum-of-contigs
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NOTE: This is a 'working draft' sequence. It currently
* consists of 21 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence
* as soon as it is available and the accession number will
* be preserved.
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1      2144: contig of 2144 bp in length
*      2145      2244: gap of unknown length
*      2245      4052: contig of 1808 bp in length
*      4053      4153: gap of unknown length
*      4153      6306: contig of 2154 bp in length
*      6307      6406: gap of unknown length
*      6407      8446: contig of 2038 bp in length
*      8445      8544: gap of unknown length
*      8545      11626: contig of 3082 bp in length
*      11627      11726: gap of unknown length
*      11727      15823: contig of 4097 bp in length
*      15824      15923: gap of unknown length
*      15924      20350: contig of 4427 bp in length
*      20351      20450: gap of unknown length
*      20451      25976: contig of 5526 bp in length
*      25977      26076: gap of unknown length
*      26077      31313: contig of 5237 bp in length
*      31314      31413: gap of unknown length
*      31414      38673: contig of 7260 bp in length
*      38674      38773: gap of unknown length
*      38774      45342: contig of 6569 bp in length
*      45343      45442: gap of unknown length
*      45443      53428: contig of 7986 bp in length
*      53429      53528: gap of unknown length
*      53529      60068: contig of 6540 bp in length
*      60069      60168: gap of unknown length
*      60169      68196: contig of 8028 bp in length
*      68197      68296: gap of unknown length
*      68297      76091: contig of 7795 bp in length
*      76092      76191: gap of unknown length
*      76192      87826: contig of 11635 bp in length
*      87827      87926: gap of unknown length
*      87927      99071: contig of 11145 bp in length
*      99072      99171: gap of unknown length
*      99172      115184: contig of 16013 bp in length
*      115185      115284: gap of unknown length
*      115285      137703: contig of 17419 bp in length
*      132704      132803: gap of unknown length
*      132804      154064: contig of 21261 bp in length
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* 154065 154164: gap of unknown length
* 154165 173556: Contig of 19392 bp in length.
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ORIGIN
Query Match 39.3%; Score 157.2; DB 12; Length 173556;
Best Local Similarity 74.6%; Pred.No. 1.2e-42;
Matches 209; Conservative 1; Mismatches 69; Indels 1; Gaps 1;

QY 81 ATGCCTGTAATCCGACACTCGGAGGCGCAAGTGGCGGATCACTGAGGTCAAGAGA 140
DB 1615 AGGCTGTATCCGACGACTTTGGGAGGCCAAGTGGCGGATCACTGAGGTCAAGAGT 1674
QY 141 TCGAGACCATCTGCGCCCAATGCTGAAACCCGCTTTACTAAATAACAAAAATAGC 200
DB 1675 TCGAGACCATCTGCGCCCAATGCTGAAACCCGCTTTACTAAATAACAAAAATAGC 1734
QY 201 TGGGCATGTGGGACACACCTGTAGTCCCACTACTCAGAGCGCGGAGATTGCACTGAGC 260
DB 1735 CAGGAGTGTGGGACACACCTGTAGTCCCACTACTCAGAGCGGAGATTGCACTGAGC 1794
QY 261 TGAGATCGACAGAGGAGCGGCAATTCACAGATCACAGAGTGAAG-CAGAGTGAAGACGCTC 319
DB 1795 CTTGATCCCGAGAGCGGAGGTTGCAATAGCCAGATGCGGCACTCGAGACTCTGTC 1854
QY 320 TCAAAAAACACACAAAAACAAAAAACCAATAGACATT 359
DB 1855 TCAAAAAATTAATAATAATAATAATAATAAAGGCACT 1894

RESULT 25
AL162739/c 84001 bp DNA linear PRI 18-MAY-2005
LOCUS
DEFINITION
Human DNA sequence from clone Rp-1155K23 on chromosome 1p31.3-32.3
Contains a Down syndrome critical region gene 5 (DSCR5) pseudogene,
a ribosomal protein S15a (RPS15A) pseudogene, a novel gene
(FLJ10884), a ribosomal protein L36 (RPL36) pseudogene and the 3'
end of a novel gene, complete sequence.
ACCESSION
AL162739
VERSION
AL162739.24 GI:21211651
KEYWORDS
HTG; DSCR5; FLJ10884; RPL36; RPS15A.
SOURCE
Homo sapiens (human)
ORGANISM
Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini;
Hominiidae; Homo.
1 (bases 1 to 84001)
REFERENCE
White,S.
Direct Submission
Submitted (13-MAY-2005) Wellcome Trust Sanger Institute, Hinxton,
Cambridgeshire, CB10 1SA, UK. E-mail enquiries: vegas@sanger.ac.uk
Clone requests: clonerequests@sanger.ac.uk
On May 25, 2002 this sequence version replaced gi:16973818.
The following abbreviations are used to associate primary accession
```

numbers given in the feature table with their source databases:

Em, EMBL; Sw, SWISSPROT; Tr, TREMBL; Mp, WORMPEP; Information on the WORMPEP database can be found at http://www.sanger.ac.uk/Projects/C_elegans/wormpep This sequence was generated from part of bacterial clone contigs of human chromosome 1, constructed by the Sanger Centre Chromosome 1 Mapping Group. Further information can be found at <http://www.sanger.ac.uk/HGP/Chri> RPS-1155K23 is from the library RPCI-5 constructed by the group of Pieter de Jong. For further details see <http://www.choi.org/bacpac/home.htm> VECTATOR: PCYPAC2

----- Genome Center
Center: Wellcome Trust Sanger Institute
Center code: SC
Web site: <http://www.sanger.ac.uk>
Contact: vega@sanger.ac.uk

This sequence was finished as follows unless otherwise noted: all regions were either double-stranded or sequenced with an alternate chemistry or covered by high quality data (i.e., phred quality >= 30); an attempt was made to resolve all sequencing problems, such as compressions and repeats; all regions were covered by at least one subclone; and the assembly was confirmed by restriction digest, except on the rare occasion of the clone being a YAC.

FEATURES

SOURCE

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terminator reads only."
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Em:AI656788.1 Em:AI700108.1 Em:AUI24205.1 Em:AUI33378.1
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CDS

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Em:BX369343.1 Em:BX382228.1 Em:BX382229.1 Em:CG988408.1
Em:CD642516.1 Em:CD656210.1 Em:CD656782.1
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complement(AL139343.9:17619..17837),
complement(AL139343.9:16064..16151),
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complement(AL139343.9:2399..2541),
complement(78805..79005),complement(68312..69714))
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repeat_region	/rpt_family="L2"
repeat_region	complement(2113. .2274)
repeat_region	/rpt_family="L2"
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repeat_region	/rpt_family="AT_rich"
repeat_region	3802. .4110
repeat_region	/rpt_family="AluY"
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repeat_region	4679. .4709
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repeat_region	8990. .9123
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repeat_region	14270. .14377
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repeats_region	17045..17462	/rpt_family="LTR45"	complement(18479..18774)	repeats_region	17045..17462	/rpt_family="LTR45"	complement(18479..18774)	repeats_region	17045..17462	/rpt_family="LTR45"	complement(18479..18774)
repeats_region	17045..17462	/rpt_family="AluSx"	19235..19484	repeats_region	17045..17462	/rpt_family="AluSx"	19235..19484	repeats_region	17045..17462	/rpt_family="AluSx"	19235..19484
repeats_region	17045..17462	/rpt_family="L1MC4a"	21809..22067	repeats_region	17045..17462	/rpt_family="L1MC4a"	21809..22067	repeats_region	17045..17462	/rpt_family="L1MC4a"	21809..22067
repeats_region	17045..17462	/rpt_family="MIR"	22107..22141	repeats_region	17045..17462	/rpt_family="MIR"	22107..22141	repeats_region	17045..17462	/rpt_family="MIR"	22107..22141
repeats_region	17045..17462	/rpt_family="(CA)n"	complement(23108..23256)	repeats_region	17045..17462	/rpt_family="(CA)n"	complement(23108..23256)	repeats_region	17045..17462	/rpt_family="(CA)n"	complement(23108..23256)
repeats_region	17045..17462	/rpt_family="MIR"	23382..23758	repeats_region	17045..17462	/rpt_family="MIR"	23382..23758	repeats_region	17045..17462	/rpt_family="MIR"	23382..23758
repeats_region	17045..17462	/rpt_family="THE1C"	complement(24728..24964)	repeats_region	17045..17462	/rpt_family="THE1C"	complement(24728..24964)	repeats_region	17045..17462	/rpt_family="THE1C"	complement(24728..24964)
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repeats_region	17045..17462	/rpt_family="MLT1C"	26691..26896	repeats_region	17045..17462	/rpt_family="MLT1C"	26691..26896	repeats_region	17045..17462	/rpt_family="MLT1C"	26691..26896
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VERSION AC068995.10 GI:1195516
KEYWORDS HTG; HTGS PHASE1; HTGS_DRAFT.
SOURCE
ORGANISM Homo sapiens (human)

Homosapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini;
Homidae; Homo.
1 (bases 1 to 195070)

REFERENCE
AUTHORS
Munoz, D.M., Adams, C., Adio-Oduola, B., Ali-ouman, F.R., Allen, C.,
Albrooks, S.L., Amaral, H.C., Are, J.R., Banks, T., Barbata, J.,
Benton, J., Bimaga, K., Blankenburg, K., Bonin, D., Bouck, J.,
Bowe, S., Brieva, M., Brown, E., Brown, N.P., Bryant, N.P., Bulay, C.,
Burke, P., Burkett, C., Burrell, K.L., Byrd, N.C., Caron, T.F.,
Carter, M., Cavazos, S.R., Chacko, J., Chavez, D., Chen, G., Chen, R.,
Chen, Z., Chowdhry, I., Christopoulos, C., Cleveland, C.D., Cox, C.,
Coyne, M.D., Dathorne, S.R., David, R., Davila, M.L., Davis, C.,
Denn, A.L., Ding, Y., Dinh, H.H., Douthwaite, K.J., Draper, H.,
Dugan-Rocha, S., Durbin, K.J., Earhart, C., Edgar, D., Edwards, C.C.,
Elhaj, C., Escoto, M., Falle, T., Ferraguto, D., Flagg, N., Ford, J.,
Foerster, P., Frantz, P., Gabler, A., Gao, J., Garcia, A., Garner, T.,
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Worley, K., Wu, C., Wu, Y., Wu, Y.F., Zhou, J., Zorrilla, S., Nelson, D.,
and Gibbs, R.
Direct Submission
Unpublished
2 (bases 1 to 195070)
Worley, K.C.
Direct Submission
Submitted (16-MAY-2000) Human Genome Sequencing Center, Department
of Molecular and Human Genetics, Baylor College of Medicine, One
Baylor Plaza, Houston, TX 77030, USA
On Dec 29, 2000 this sequence version replaced gi:11024742.
----- Genome Center
Center: Baylor College of Medicine
Center code: BCM
Web site: http://www.hgsc.bcm.tmc.edu/
Contact: hgsc-help@bcm.tmc.edu
----- Project Information
Center project name: HBGH
Center clone name: RP11-463G10
----- Summary Statistics
Sequencing vector: M13, L08821
Chemistry: Dye-terminator Big Dye: 100% of reads
Assembly program: Phrap, version 0.990329
Consensus quality: 160818 bases at least Q40
Consensus quality: 177694 bases at least Q30
Consensus quality: 184433 bases at least Q20
Estimated insert size: 185987; sum-of-contigs estimation
Quality coverage: 0x in Q20 bases; agarose-1p estimation

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32637. 32736
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78735. 78834

Quality coverage: 3.4x in Q20 bases; sum-of-contigs estimation

* NOTE: Estimated insert size may differ from sequence length
* (see http://www.hgsc.bcm.tmc.edu/docs/genbank_draft_data.html).
* NOTE: This is a 'working draft' sequence. It currently
* consists of 23 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence
* as soon as it is available and the accession number will
* be preserved.

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20258: gap of unknown length
32636: contig of 12378 bp in length
32736: gap of unknown length
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49756: gap of unknown length
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108563: contig of 13210 bp in length
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120335: contig of 11672 bp in length
120435: gap of unknown length
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155046: contig of 7609 bp in length
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162104: contig of 658 bp in length
162204: gap of unknown length
168086: contig of 5882 bp in length
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175732: contig of 7546 bp in length
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180404: contig of 4572 bp in length
180504: gap of unknown length
182399: contig of 1895 bp in length
182400: gap of unknown length
184812: contig of 2313 bp in length
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187927: contig of 3015 bp in length
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189826: contig of 1799 bp in length
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192637: contig of 2711 bp in length
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ORIGIN

Query Match 39.2%; Score 156.8; DB 12; Length 195070;
 Best Local Similarity 71.0%; Pred. No. 1.8e-42;
 Matches 206; Conservative 1; Mismatches 83; Indels 0; Gaps 0;

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Oy      AGGCCAGCATGACCTGCTGATGATCCCTGATCCAGCACTTCGGAGGCCAGAGTGG 117
          |||||
Db      2493 AGCCCTGGCCAGCGCGGTACTCATGCTCTGTAATCCAGCACTTGGAGGTGAGCGG 2434
          |||||
Oy      118 GCGGATCACTGAGGTCAAGAGATGAGACCATCTGCGCAACATGTTGAAACCCCGTCT 177
          |||||
Db      2433 GTGATCACTGATGATCAGAGTTCAAGACCGCTGGCAACATGACCAAAACCCGCT 2374
          |||||
Oy      178 TTACTTAAATATCAAAAAATAGCTGGCATGTGGCAACACCTTATGCTCCAGTACTC 237
          |||||
Db      2373 CTACTTAAATATCAAAAAATAGCTGGCATGTGGCAACCTTATGCTCCAGTACTC 2314
          |||||
Oy      238 AGGAGCCGAGATTCAGTGAAGTCAAGATCGCAGAGTGAGCCGAATATCAGATACAGA 297
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Db      2313 AGGAGGCTGAGGAGGAGAAATTACAGTGGCGGAGATCGACCACTAGACCTCCAGCTG 2254
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Oy      2253 GTGAGCAGAGTGAGACCCGCTCTCAAAAAACAACAACAAAAAA 347
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RESULT 28
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 LOCUS Homo sapiens chromosome 1 clone RP4-633K13 map p34.3-36.11, 9
 DEFINITION unordered pieces.
 ACCESSION AL353145
 VERSION AL353145.4 GI:9796994
 KEYWORDS HTG; HTGS_PHASE1; HTGS_CANCELLED.
 SOURCE Homo sapiens (human)
 ORGANISM Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

REFERENCE

Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini;
 Homiidae; Homo.
 1
 McIay, K.
 Direct Submission
 Submitted (09-JUL-2001) Sanger Centre, Hinxton, Cambridgeshire,
 CB10 1SA, UK. E-mail enquiries: humquerry@sanger.ac.uk
 requests: clonerequest@sanger.ac.uk
 On Aug 12, 2000 this sequence version replaced gi:9213061.

 Genome Center
 Center: Sanger Centre
 Center code: SC
 Web site: http://www.sanger.ac.uk
 Contact: humquerry@sanger.ac.uk

 Project Information
 Center project name: d0633K13

 Summary Statistics
 Assembly program: XGAP4; version 4.5
 Sequencing vector: plasmid; 108752; 100% of reads
 Chemistry: Dye-terminator ABI; 1% of reads
 Chemistry: Dye-terminator Big Dye; 74% of reads
 Chemistry: Dye-terminator ET-amersham; 24% of reads
 Consensus quality: 107404 bases at least Q40
 Consensus quality: 10616 bases at least Q20
 Insert size: 110572; sum-of-contigs
 Insert size: 124958; 7.7% error; agrose-fp
 Quality coverage: 3.70x in Q20 bases; sum-of-contigs Quality
 coverage: 3.47x in Q20 bases; agrose-fp

COMMENT

* NOTE: This is a 'working draft' sequence. It currently
 * consists of 9 contigs. The true order of the pieces
 * is not known and their order in this sequence record is
 * arbitrary. Gaps between the contigs are represented as
 * runs of N, but the exact sizes of the gaps are unknown.
 * This record will be updated with the finished sequence
 * as soon as it is available and the accession number will
 * be preserved.

FEATURES

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misc_feature

misc_feature

misc_feature

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Query Match      39.2% Score 156.6; DB 12; Length 111372;
Best Local Similarity 71.1%; Pred.No.1.6e-42;
Matches 207; Conservative 0; Mismatches 84; Indels 0; Gaps 0;
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QY      141 TCGAGACCATCTGGCCCAACATGTGAAACCCGCTTTACTTAAATAACAAAAATAGC 200
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QY      201 TGGGATGATGGCAACACCTGTACTCCAGCTACTCAGAGACCGGAGATTCAGTGAAC 260
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QY      261 TGAGATCCAGAGTGAAGCCGAATATCAGATCAGATGAGAGAGAGAGAGAGAGAGAGAG 320
DB      104604 CTTGACCCCAAGAGTTGAGAGTGAAGTGAAGTGAAGTGAAGTGAAGTGAAGTGAAGT 104545
QY      321 CAAAAACAACAACAAAAACAAAAACCAATAGACATTTGTCATCTCGCG 371
DB      104544 AATATCAAAACAAAAACAAAAACAAATATATTAAGATTAAGTGTGG 104494
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LOCUS      AC022252
DEFINITION Homo sapiens clone RP11-28012, WORKING DRAFT SEQUENCE, 28 unordered
ACCESSION AC022252
VERSION    AC022252.2 GI:7249106
KEYWORDS   HTG; HTGS_PHASE1; HTGS_DRAFT.
SOURCE     Homo sapiens (human)
ORGANISM   Homo sapiens
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            Homo sapiens, clone RP11-28012
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            Biren, B., Linton, L., Nusbaum, C., Lander, E., Abraham, H., Allen, N.,
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            Norman, C. H., O'Connor, T., O'Donnell, P., Olivar, T. M., Peterson, K.,

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TITLE JOURNAL COMMENT

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Pierre, N., Pisan, C., Pollara, V., Raymond, C., Riley, R., Rotman, D.,
Roy, A., Santos, R., Severy, P., Spencer, B., Scange-Thomann, N.,
Stojanovic, N., Subramanian, A., Talamas, J., Testa, S., Theodore, J.,
Tirrell, A., Vassiliev, H., Viel, R., Vo, A., Wu, X., Wyman, D., Ye, W. J.,
Zimmer, A. and Zody, M.
Direct Submission
Submitted (27-JUN-2000) Whitehead Institute/MIT Center for Genome
Research, 320 Charles Street, Cambridge, MA 02141, USA
On Mar 16, 2000 this sequence version replaced gi:6778512.
All repeats were identified using RepeatMasker:
Smit, A.F.A. & Green, P. (1996-1997)
http://ftp.genome.washington.edu/RM/RepeatMasker.html
-----
Genome Center
Center: Whitehead Institute/ MIT Center for Genome Research
Center code: WIBR
Web site: http://www-seq.wi.mit.edu
Contact: sequence.submissions@genome.wi.mit.edu
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Project Information
Center project name: L4795
Center clone name: 28_012
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Summary Statistics
Sequencing vector: M13; M77815; 100% of reads
Chemistry: Dye-terminator Big Dye; 100% of reads
Assembly program: Phrap; version 0.960731
Consensus quality: 132159 bases at least Q40
Consensus quality: 141146 bases at least Q30
Consensus quality: 144793 bases at least Q20
Insert size: 157000; agarose-fp
Insert size: 148234; sum-of-coverage
Quality coverage: 3.1 in Q20 bases; agarose-fp
Quality coverage: 3.2 in Q20 bases; sum-of-coverage
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* NOTE: This is a 'working draft' sequence. It currently
* consists of 28 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence
* as soon as it is available and the accession number will
* be preserved.
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*      1564      1663: gap of 100 bp
*      1664      2811: contig of 1148 bp in length
*      2812      2911: gap of 100 bp
*      2912      4293: contig of 1382 bp in length
*      4294      4393: gap of 100 bp
*      4394      4577: contig of 364 bp in length
*      4578      4857: gap of 100 bp
*      4858      6513: contig of 1656 bp in length
*      6514      6613: gap of 100 bp
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*      8469      8569: gap of 100 bp
*      8569      10457: contig of 1889 bp in length
*      10458      10557: gap of 100 bp
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*      16827      20428: contig of 3602 bp in length
*      20429      20528: gap of 100 bp
*      20529      24488: contig of 3960 bp in length
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*      28868      28967: gap of 100 bp
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*      33308      33407: gap of 100 bp
*      33408      35317: contig of 1910 bp in length
*      35318      35417: gap of 100 bp
*      35418      39417: contig of 4000 bp in length
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*      39518      45499: contig of 5982 bp in length

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* 69807 69906: gap of 100 bp
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* 77967 78097 78096: gap of 100 bp
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* 88133 88232: gap of 100 bp
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/feature="assembly_fragment"
1564. .1663
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/misc_feature
/feature="assembly_fragment"
1664. .2811
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2812. .2911
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/misc_feature
/feature="assembly_fragment"
2912. .4293
/feature="assembly_fragment"
4294. .4393
/estimated_length=100
/misc_feature
/feature="assembly_fragment"
4394. .4757
/feature="assembly_fragment"
clone end: r7
vector_side:right"
4758. .4857
/estimated_length=100
/misc_feature
/feature="assembly_fragment"
4858. .6513
/feature="assembly_fragment"
6514. .6613
/estimated_length=100
/misc_feature
/feature="assembly_fragment"
6614. .8468
/feature="assembly_fragment"
8469. .8568
/estimated_length=100
/misc_feature
/feature="assembly_fragment"
8569. .10457
/feature="assembly_fragment"
10458. .10557
/estimated_length=100
/misc_feature
/feature="assembly_fragment"
10558. .12036
/feature="assembly_fragment"
12037. .12136
/estimated_length=100
/misc_feature
/feature="assembly_fragment"
12137. .13936
/feature="assembly_fragment"
13937. .14036
/estimated_length=100
/misc_feature
/feature="assembly_fragment"
14037. .16726
/feature="assembly_fragment"
16727. .16826
/estimated_length=100
/misc_feature
/feature="assembly_fragment"
16827. .20428
/feature="assembly_fragment"

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gap 20429. .20528
/misc_feature /estimated_length=100
20529. .24168
/feature="assembly_fragment"
gap 24489. .24588
/misc_feature /estimated_length=100
24589. .28867
/feature="assembly_fragment"
gap 28868. .28967
/misc_feature /estimated_length=100
28968. .33307
/feature="assembly_fragment"
gap 33308. .33407
/misc_feature /estimated_length=100
33408. .35317
/feature="assembly_fragment"
/misc_feature /note="assembly_fragment"
clone_end:SP6
vector_side:left"
35318. .35417
/misc_feature /estimated_length=100
35418. .39417
/feature="assembly_fragment"
gap 39418. .39517
/misc_feature /estimated_length=100
39518. .45499
/feature="assembly_fragment"

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Query Match 39.2%; Score 156.6; DB 12; Length 150934;
Best Local Similarity 73.1%; Pred. No. 1,8e-42;
Matches 201; Conservative 0; Mismatches 74; Indels 0; Gaps 0;

QY 81 ATGCGTGAATCCAGAGACTTCGGAGAGCCCAAGTGGCCGATCACCTGAGTCAAGAGA 140
Db 87071 ATGCGTGAATCCAGAGACTTTGGAGAGCCCAAGCAGGCCGATCAGTCAAGAGT 87130
QY 141 TCGAGACCATCTCTGGCCACATGCTGAAACCCGCTTTTACTAAATAACAAAAATAGC 200
Db 87131 TCAAGACCAAGCTGACCAACATGCTGAAACCCGCTCTACTAAATAACAAAAATAGC 87190
QY 201 TGGGCATGTGTGGCACAACCTGTAGTCCAGCTACTAGAGAGCCGAGATTGACAGTACG 260
Db 87191 TGGGTGTGTGGCACAAGCCTGTAGTCCAGCTACTAGAGAGCTGAGATGGAATCT 87250
QY 261 TGAGATCGACAGAGTGAACCCGAATCACAGATCACAGATGAGAGTGAAGACKCGTCT 320
Db 87251 CTTGAACCCGGAGAGTGAAGATCTGACTGAGCCGAGACTCGCCTGCTGACAGAGCG 87310
QY 321 CAAAAACACACAAAAACAAAAAACATTAAGA 355
Db 87311 AGACACCATCTCAAAAAAAAAAAAAAAAAAGGA 87345

RESULT 30
AC108040/c 163521 bp DNA 1linear HTG 30-JAN-2002
LOCUS Homo sapiens chromosome 4 clone RP11-210010, WORKING DRAFT
DEFINITION SEQUENCE, 4 unordered pieces.
ACCESSION AC108040.2 GI:18425316
VERSION HTG; HTGS PHASE1; HTGS_DRAFT; HTGS_FULLTOP.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini;
Hominae; Homo.
1 (bases 1 to 163521)
REFERENCE 1
AUTHORS Waterston,R.H.
TITLE The sequence of Homo sapiens clone
JOURNAL Unpublished
REFERENCE 2 (bases 1 to 163521)
AUTHORS Waterston,R.H.
TITLE Direct Submission
JOURNAL Submitted (24-JAN-2002) Genome Sequencing Center, Washington

```

University School of Medicine, 4444 Forest Park Parkway, St. Louis,
MO 63108, USA
On Jan 30, 2002 this sequence version replaced gi:18308830.

----- Genome Center -----
Center: Washington University Genome Sequencing Center
Center code: WUGSC
Web site: <http://genome.wustl.edu/gsc/index.shtml>
Contact: submissions@wustl.edu
----- Project Information -----
Center project name: H_NH0210010

----- Summary Statistics -----
Sequencing vector: M13; 0%
Sequencing vector: Plasmid; 100%
Chemistry: Dye-terminator Big Dye; 100% of reads
Chemistry: Dye-terminator Big Dye; 100% of reads
Assembly program: Phrap; version 0.990319
Consensus quality: 162304 bases at least Q40
Consensus quality: 162444 bases at least Q30
Consensus quality: 162529 bases at least Q20
Insert size: 189000; agarose-fp
Insert size: 163221; sum-of-contrigs
Quality coverage: 8.88 in Q20 bases; sum-of-contrigs
Quality coverage: 8.85 in Q20 bases; sum-of-contrigs

* NOTE: This is a 'working draft' sequence. It currently
* consists of 4 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence
* as soon as it is available and the accession number will
* be preserved.

1 1564: contig of 1564 bp in length
* 1565 1664: gap of unknown length
* 1665 88984: contig of 87320 bp in length
* 88985 89084: gap of unknown length
* 89085 112702: contig of 23618 bp in length
* 112703 112802: gap of unknown length
* 112803 163521: contig of 50719 bp in length.
Location/Qualifiers

FEATURES
source
1. 163521
/organism="Homo sapiens"
/mol_type="genomic DNA"
/db_xref="taxon:9606"
/chromosome="4"
/clone="RP11-210010"
1. 1564
/note="assembly_name:Contig25
clone_end:SP6
vector_side:left"
1565. 1664
/estimated_length=unknown
1665. 88984
/note="assembly_name:Contig28
clone_end:T7
vector_side:left"
88985. 89084
/estimated_length=unknown
89085. 112702
/note="assembly_name:Contig26"
112703. 112802
/estimated_length=unknown
112803. 163521
/note="assembly_name:Contig27"

ORIGIN

Query Match 39.2%; Score 156.6; DB 12; Length 163521;
Best Local Similarity 71.4%; Pred. No. 1.9e-42;
Matches 222; Conservative 1; Mismatches 80; Indels 8; Gaps 1;
37 AATATTATTAAGACATTGTGAGCCAGGACATGACACTGGCTGATGCTTATATCCAG 96

Db 98071 AAAAAAAAAAATACAGAGCTGCTGGGTGTATGGCTATGCTTTATCTTAC 98012
Qy 97 CACTTGGGAGCCAGAGTGGCGGATCAGCTGAGGTCAAGATGAGACCATCTGGC 156
Db 98011 CACTTGGGAGCCAGAGTGGCGGATCAGCTGAGGTCAAGATGAGACCATCTGGC 97952
Qy 157 CAACATGGTGAACCCCGCTTTCTAATAAATATCAAAATATGCTGGGCATGTCGACA 216
Db 97951 CAACATGGTGAACCCCGCTTTCTAATAAATATCAAAATATGCTGGGCATGTCG 97892
Qy 217 CACCTGTAGTCCAGCTACTCAGAGCCGAGATTGACGTGAGCTGAGATGCAAGTGA 276
Db 97891 CGCTTGTATATCCAGCTACTCAGAGCCGAGATTGACGTGAGCTGAGATGCA 97840
Qy 277 GCCAATACAGATCAAGAGTGAAGAGTGAAGCCGCTCTCAAAACACACAA 336
Db 97839 TGCCATTGCACTTGGCGCTGGAGACAAAGTGAACCTCAATCAAAAAATTAACA 97780
Qy 337 AATCAAAAAA 347
Db 97779 AATTAATACA 97769

RESULT 31

LOCUS AC114480 196773 bp DNA linear PRI 29-JAN-2003
DEFINITION Homo sapiens chromosome 3 clone RP11-755B10, complete sequence.
AC114480
AC114480.2 GI:28014590
VERSION HTG.
KEYWORDS
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens

REFERENCE
AUTHORS Kaul,R.K., Olson,M.V., Zhou,Y., James,R.A., Rouse,G., Wu,Z.,
Saenphimachak,C., Buckley,D., Kibukawa,M., Raymond,C. and
Haugen,E.D.
1 (bases 1 to 196773)
2 (bases 1 to 196773)
Direct Submission
Unpublished

TITLE
JOURNAL
REFERENCE
AUTHORS Kaul,R.K., Olson,M.V., Raymond,C. and Haugen,E.D.
Direct Submission
Submitted (09-MAR-2002) Genome Center, University of Washington,
Box 352145, Seattle, WA 98195, USA
3 (bases 1 to 196773)
Kaul,R.K., Olson,M.V., Zhou,Y., James,R.A., Rouse,G., Wu,Z.,
Saenphimachak,C., Buckley,D., Kibukawa,M., Raymond,C. and
Haugen,E.D.
Direct Submission
Submitted (29-JAN-2003) Genome Center, University of Washington,
Box 352145, Seattle, WA 98195, USA
On Jan 29, 2003 this sequence version replaced gi:19310299.

TITLE
JOURNAL
COMMENT
On Jan 29, 2003 this sequence version replaced gi:19310299.

----- Genome Center -----
Center: University of Washington Genome Center
Center Code: WUGC
Web site: <http://www.genome.washington.edu>
Contact: wugc@u.washington.edu
----- Project Information -----
Center project name: chr-3
Center clone name: RP11-755B10 (bc0671)
----- Summary Statistics -----
Sequencing vector: unknown; 3% of reads
Sequencing vector: plasmid; 97% of reads
Chemistry: Dye-terminator ET; 94% of reads
Chemistry: Dye-terminator Big Dye; 6% of reads
Assembly program: Phrap; version 0.990319
Consensus quality: 196438 bases at least Q40
Consensus quality: 196773 bases at least Q30
Consensus quality: 196773 bases at least Q20
Insert size: 196773; sum-of-contrigs

Quality coverage: 8.4x in Q20 bases; sum-of-contigs

Overlapping Sequences:

5: RP11-229A12 (UMGC:bc0315) AC092418, 11308-bp overlap
3: RP11-680P23 (UMGC:bc0527) AC099558, 118740-bp overlap

Sequence Quality Assessment:

This entry has been annotated with sequence quality estimates computed by the Phrap assembly program. All manually edited bases have been reduced to quality zero. Quality levels above 40 are expected to have less than 1 error in 10,000 bp. Base-by-base quality values are not generally visible from the GenBank flat file format but are available as part of this entry's ASN.1 file.

This sequence was finished as follows unless otherwise noted: all regions were either double-stranded or sequenced with an alternate chemistry or covered by high quality data (i.e., Phred quality >= 30); an attempt was made to resolve all sequencing problems, such as compressions and repeats; all regions were covered by at least one plasmid subclone or more than one M13 subclone; and the assembly was confirmed by restriction digest.

Sequence Validation:

This sequence has been validated by Multiple Complete Digest fingerprinting. Comparison of the experimentally derived digest fragments with sequence-predicted fragments is given below. The electronically-digested sequence consists of both insert and vector, in order to accurately represent the entire circular BAC. Small fragments below a variable cutoff (approximately 400-800 bp) are not resolved in the fingerprint and hence do not appear in the table. There are no significant remaining discrepancies between the experimental and predicted values. Uniquely ordered fragments are separated by dashed lines.

ECORI

HindIII

BglII

SeqDerMap	FingerPrint	SeqDerMap	FingerPrint	SeqDerMap	FingerPrint
8696	8856	3845	3999	10997	10643
6	<800	6382	6490	2067	2051
4995	4998	512	<800	12782	12807
2246	2241	449	<800	13590	12807
4061	4055	11421	11078	13168	12807
2665	2694	3245	3293	3447	3526
519	<800	3752	3826	5493	5591
1442	1386	1035	1032	1619	1545
6438	6457	881	873	9870	9617
2522	2531	5764	5726	6707	6693
9191	9289	6793	6788	3014	3167
2510	2531	889	873	4466	4431
138	<800	740	<800	16871	17658
2876	2910	2548	2558	1067	1017
1971	1982	2797	2775	5594	5591
812	796	5345	5298	10765	10643

1196	1166	552	<800	6602	6693
3348	3354	540	<800	4314	4431
11391	11222	1886	1862	846	836
16746	16576	5371	5298	3702	3743
340	<800	12306	11906	2801	2913
2626	2694	2938	2906	266	<800
4361	4390	321	<800	5324	5342
437	<800	14621	14206	3164	3280
2459	2531	7088	7077	7697	7733
18703	18633	684	<800	2197	2308
2240	2241	4160	4387	1501	1449
14113	13849	657	<800	1359	1311
27	<800	677	<800	15679	16033
1993	1982	669	<800	5370	5342
28749	28884	4437	4387	10381	10125
1970	1982	3581	3651	12752	12807
15338	15091	1088	1032		
417	<800	6707	6788		
21440	21694	6490	6490		
328	<800	4499	4387		
4697	4783	2212	2210		
478	<800	17962	17884		
987	967	4016	4136		
		2063	2063		
		8874	8921		
		3328	3518		
		22128	22660		
		7472	7554		
		1747	1718		

FEATURES

source

Location/Qualifiers
1..196773
/organism="Homo sapiens"
/mol_type="genomic DNA"
/db_xref="taxon:9606"
/chromosome="13"
/clone="RP11-755B10"
/clone_id="RP11 human BAC library 11"

ORIGIN

Query Match 39.2%; Score 156.6; DB 5; Length 196773;
Best Local Similarity 71.4%; Pred. No. 2.1e-42;
Matches 222; Conservative 1; Mismatches 80; Indels 8; Gaps 1;
37 AATATTAAATAGACATTGTCAGCCAGCATGACACTGGCTGAATGCTGTAATCCAG 96

```

Db      31090 AAAAAATATATCAAAAAATACAGAAAGCTGGCTGCGGTGATGGCTTCATGCTTGTATATCTTAG 31144
Oy      97 CACTTCGGGAGGCCAAGGTGGCGGATCACCTGAGTCAAGAGATGAGACCATCTGGC 156
Db      31150 CACTTTGGGAGGCCAAGGCAAGTGGATGATCACTGAGGTCAAGAGTTGCAAGCTGAGCTGGC 31209
Oy      157 CAACATGTTGAAACCCCGTCTTTTCTTAAAAATACAAAAAATAGTGGGCAATGGTGGCA 216
Db      31210 CAACATGTTGAAACCCCGTCTTCTTAAAAATACAAACATTACTGGGCTTGGTGGCGG 31265
Oy      217 CACCTGTATGTCACAGTACTCAGGAGCGGAGATTGCAGTAGCTGATCGAGAGTGA 276
Db      31270 CGCTTGTATATCCAGCTACTGAGGAGGCAAGGTTGCAAGTGAAGCTTGATC-----A 31322
Oy      277 GCCGAAATTCACAGATCAACAGATGAGCAGAGTGAAGACCCGCTCAAAAACAAACA 336
Db      31322 TGCCATGTCACACTGCGCTGGGAGACAAGAGTGAAATCTCATCTCAAAAAAATAAACA 31382
Oy      337 AAACAAAAAA 347
Db      31382 AATAAATACA 31392

RESULT 32
AC016552/c
LOCUS      102008 bp      DNA      linear      HTG 19-APR-2001
DEFINITION Homo sapiens chromosome 5 clone CTC-285M15, WORKING DRAFT SEQUENCE,
4 ordered pieces.
AC016552
AC016552.5 GI:7711562
HTG: HTGS_PHASE2; HTGS_DRAFT; HTGS_ACTIVEPIN.
KEYWORDS   Homo sapiens (human)
ORGANISM   Homo sapiens
            Eumaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
            Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini;
            Homiidae; Homo.
REFERENCE   1 (bases 1 to 102008)
            DOE Joint Genome Institute.
            Sequencing of Human Chromosome 5
            Unpublished
            2 (bases 1 to 102008)
            DOE Joint Genome Institute.
            Direct Submision
            Submitted (04-DEC-1999) Production Sequencing Facility, DOE Joint
            Genome Institute, 2800 Mitchell Drive, Walnut Creek, CA 94598, USA
            On May 6, 2000 this sequence version replaced gi:7710241.
COMMENT
-----Genome Center
Center: Joint Genome Institute
Center Code: JGI
Web site: http://www.jgi.doe.gov
-----
Project Information
Center Project Name: 294932
Center clone name: CIT-HSFC_285M15
-----
Summary Statistics
Consensus quality: 92975 bases at least Q40
Consensus quality: 98251 bases at least Q30
Consensus quality: 100181 bases at least Q20
Estimated insert size: 16000; pulse field gel estimation
Estimated insert size: 10185; sum-of-contigs estimation
Quality coverage: 5.46 in Q20 bases; pulse field gel estimation
Quality coverage: 5.69 in Q20 bases; sum-of-contigs estimation.
NOTE: This is a 'working draft' sequence. It currently
* consists of 4 contigs. Gaps between the contigs
* are represented as runs of N. The order of the pieces
* is believed to be correct as given, however the sizes
* of the gaps between them are based on estimates that have
* provided by the submittor.
* This sequence will be replaced
* by the finished sequence as soon as it is available and
* the accession number will be preserved.

```

FEATURES	source	Location/Qualifiers
*	1	63901: contig of 63901 bp in length
*	63902	64001: gap of unknown length
*	64002	92415: contig of 28414 bp in length
*	92416	92515: gap of unknown length
*	92516	101204: contig of 8689 bp in length
*	101205	101304: gap of unknown length
*	101305	102008: contig of 704 bp in length.
ORIGIN		
gap		1..102008 /organism="Homo sapiens"
gap		/mol_type="genomic DNA"
gap		/db_xref="taxon:9606"
gap		/chromosome="5"
gap		/clone="CTC-285M15"
gap		/clone_id="Caltech human BAC library C"
gap		63902..64001 /estimated_length=unknown
gap		92416..92515 /estimated_length=unknown
gap		101205..101304 /estimated_length=unknown
gap		/estimated_length=unknown
Query Match	39.1%	Score 156.4; DB 12; Length 102008;
Best Local Similarity	76.6%	Pred. No. 1.8e-42; Indels 3; Gaps 2;
Matches 216; Conservative 1; Mismatches 62;		
Db	81	ATGCTGTAACTCCAGCACTTCGGAGGCGCAAGGTGGCGGATCACTGTAGTCAAGAGA 140
QY	71686	ATGCTGTAACTCCAGCACTTCGGAGGCGCAAGGTGGCGGATCACTGTAGTCAAGAGA 7162
Db	141	TCGAGACCATTCTTGCCCAATGCTGAAACCCGCTCTTACTTAAATAATACAAAAATAGC 200
QY	71626	TCGAGACCATTCTTGCCCAATGCTGAAACCCGCTCTTACTTAAATAATACAAAAATAGC 7156
Db	201	TGGGCATGGT--GGCACAACCTGTATCCAGTACTCAAGAGCCGAGATTCGAGTGA 258
QY	71566	CAGGTGTGTGTGGGACACCGCTGTATCCAGTACTCAAGAGCTGAGGCGAGGCGAGCAAT 7150
Db	259	GCTGAGATTCGAGAGTGAAGCCGAATATCAGATCAAGATCGAGAGTGAAGCKCCGT 318
QY	71506	CGCTTGAACCCAGGAGGTGAAGCCTGTGATCCAGTCTCGAGCGTG--GGCAACAGAGTGAAGCTCGT 7144
Db	319	CTCAAAAACAAACAACAAAAAACAAAAAACCATTAAGCATTG 360
QY	71447	CTCAAAAACAAACAACAAAAAACAAAAAACCATTAAGTGCCTTG 71406
RESULT 33		
AC008499/c	244525 bp	DNA linear PRI 26-JAN-2002
LOCUS		
DEFINITION		Homo sapiens chromosome 5 clone CTC-43803, complete sequence.
ACCESSION		AC008499
VERSION		AC008499.8 GI:18376856
KEYWORDS		HTG.
SOURCE		Homo sapiens (human)
ORGANISM		Homo sapiens
REFERENCE		Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
AUTHORS		Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini;
TITLE		Hominidae; Homo
JOURNAL		1 (bases 1 to 244525)
REFERENCE		DOE Joint Genome Institute and Stanford Human Genome Center.
AUTHORS		Direct Submission
JOURNAL		2 (bases 1 to 244525)
REFERENCE		DOE Joint Genome Institute.
AUTHORS		Direct Submission
TITLE		Submitted (06-DEC-2001) DOE Joint Genome Institute, 2800 Mitchell

REFERENCE
1 (pages 1 to 18725)
AUTHORS
Antonelli, A., Bass, D., Benjamin, B., Bera, J., Blakeley, R. W.,
Bouffard, G. G., Brinkley, C., Brooks, S., Chu, G., Coleman, H.,
Franks, S., Fukenko, T., Gestole, M., Greene, A., Guan, X., Gupta, J.,

Hunter, G., Hurler, B., Idol, J. R., Kwong, P., Lalic, P., Larson, S., Lee-Lin, S.-Q., Legaspi, R., Madden, M., Maduro, Q. L., Maduro, V. B., Margulies, E. H., Masiello, C., Maskeri, B., McDowell, J., Montemayor, C., Mullikin, J. C., Park, M., Prasad, A., Puri, O., Rant, K., Reddik-Dugue, N., Sante, A., Schandler, K., Schueler, M. G., Sison, C., Stantrop, S., Tave, A., Thomas, J. W., Thomas, P. J., Tsipouri, V., Ung, L., Vogt, J. L., Wetherby, K. D., Withers, T. R., Young, A. and Green, E. D.
 NISC Comparative Sequencing Initiative
 Unpublished
 2 (bases 1 to 182725)
 Green, E. D.
 Direct Submission
 Submitted (23-NOV-2005) NIH Intramural Sequencing Center, 5625 Fishers Lane, Rockville, MD 20852, USA
 3 (bases 1 to 182725)
 Green, E. D.
 Direct Submission
 Submitted (31-JAN-2006) NIH Intramural Sequencing Center, 5625 Fishers Lane, Rockville, MD 20852, USA
 On Jan 31, 2006 this sequence version replaced gi:82617757.

 Genome Center
 Center: NIH Intramural Sequencing Center
 Center code: NISC
 Web site: <http://www.nisc.nih.gov>
 Contact: nisc_zoo@ngri.nih.gov

 Project Information
 Center project name: mrn
 Center clone name: 041K03

The sequence data in this record represents an 'enhanced' version of a Phase 2 submission. Specifically, the indicated order and orientation of each sequence contig has been established using one or more of the following: read-pair data from individual subclones, overlaps with neighboring clones, alignment with available reference sequence (e.g., human), and/or confirmation by PCR testing. In addition, the sequence assembly is generally based on at least 8X average coverage in Q20 bases and has been reviewed to rule out gross misassemblies, the low-quality ends of sequence contigs have been trimmed away, and each base is associated with a Phrap-derived quality score.

 Summary Statistics
 Sequencing vector: plasmid; n/a; 100% of reads
 Chemistry: Dye-terminator Big Dye; 100% of reads
 Assembly program: Phrap; version 0.990319
 Consensus quality: 179767 bases at least Q40
 Consensus quality: 180990 bases at least Q30
 Consensus quality: 181586 bases at least Q20
 Insert size: 214000; agarose-fp
 Insert size: 181925; sum-of-contigs
 Quality coverage: 9.44x in Q20 bases; agarose-fp
 Quality coverage: 11.11x in Q20 bases; sum-of-contigs

 NOTE: This is a 'working draft' sequence. It currently consists of 9 contigs. Gaps between the contigs are represented as runs of N. The order of the pieces is believed to be correct as given, however the sizes of the gaps between them are based on estimates that have been provided by the submitter.
 * This sequence will be replaced
 * by the finished sequence as soon as it is available and
 * the accession number will be preserved.

* 22544 22543: contig of 22543 bp in length
 * 22644 22643: gap of unknown length
 * 31875 31875: contig of 9232 bp in length
 * 31876 31875: gap of unknown length
 * 122998 122998: contig of 91023 bp in length
 * 123099 123098: gap of unknown length
 * 123099 123098: contig of 13389 bp in length
 * 136488 136487: gap of unknown length
 * 136588 136587: gap of unknown length
 * 139369 139369: contig of 2782 bp in length

* 139370 139469: gap of unknown length
 * 139470 148608: contig of 9139 bp in length
 * 148609 148708: gap of unknown length
 * 148709 157275: contig of 8567 bp in length
 * 157276 157375: gap of unknown length
 * 157376 179801: contig of 22426 bp in length
 * 179802 179901: gap of unknown length
 * 179902 182725: contig of 2824 bp in length.

FEATURES
 source
 1..182725
 /organism="Callitriche moloch"
 /mol_type="genomic DNA"
 /db_xref="taxon:9523"
 /clone_lib="LB5-41K3"
 /clone_lib="LB5"
 /note="BAC resource: <http://bacpac.chori.org/>"
 1..22543
 /note="assembly_fragment
 missing T7 clone end on 5' end of insert"
 1..18769
 /note="clone overlaps with GenBank Accession Number AC174418 clone LB5-285122 (center project name mrn)"
 22544..22643
 /estimated_length=unknown
 22644..31875
 /note="assembly_fragment"
 31876..31975
 /estimated_length=unknown
 31976..122998
 /note="assembly_fragment"
 50743..182725
 /note="clone overlaps with GenBank Accession Number AC172795 clone LB5-29099 (center project name mro)"
 122999..123098
 /estimated_length=unknown
 123099..136487
 /note="assembly_fragment"
 136488..136587
 /estimated_length=unknown
 136588..139369
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 139370..139469
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 139470..148608
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 148609..148708
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 148709..157275
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 157276..157375
 /estimated_length=unknown
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ORIGIN
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 Best Local Similarity 70.5%; Pred. No. 3.9e-42;
 Matches 225; Conservative 1; Mismatches 83; Indels 10; Gaps 1;

QY 81 ATGCTGTATCCAGCAGCTTGGAGGCGCAAGTGGCGGATCACTGAGGTCAAGGA 140
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 DB 23508 ACGCTGTATCCAGCAGCTTGGAGGCGTGAATGCTGAGGTCAAGGT 23449
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 QY 141 TCGAGCATCTGGCCCAACATGGTGAACCCCGCTTTACTTAATAAATACAAATAATGC 200
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 DB 23448 TAGAGACCAAGCTGGCCCAACATGGTGAACCCCGCTTTACTTAATAAATAAGAAATTAGC 23389

OY		201	TGGCAGTGTGGCACACACCCTGATGCCCGACTACTGAGGCGGAGATTGCAGTGACC	260
DB	23388	TGGCAGTGTGGCACACACCCTGATGCCCGACTACTGAGGCGGAGATTGCAGTGACC	23329	
OY		261	TGAGATCGCAGAGTGAGCCGGAATCATCAGATTCACAGAGTAGCAGATGAGACKCCGTCT	320
DB	23328	CGAGATCA-----CAACTGCATCTCCAGCCCTGGGAAAGAGGGAGACTTCATCT	23279	
OY		321	CAAAAACAACAACAAAAAACAATAACATTAAGACATTTGTCATTCGCGTTCCAGAC	380
DB	23278	CAAAAACAACAAAAAACAAGAATTAAGTGGCAGTAGATGGCTCAGCGCTTAATCCATCAC	23219	
OY		381	TATTGCAGGAGACCAAAAA 399	
DB	23218	TTTGAAAGGCCAACGAGAA 23200		
RESULT_36	DJ534K4			
LOCUS	DJ534K4	216387 bp	DNA	linear PRI 23-DEC-1998
DEFINITION	Homo sapiens S1c4 gene, partial cds; psi and hypothetical protein genes, complete cds; and S171 gene, partial cds.			
ACCESSION	AF109907			
VERSION	AF109907.1	GI:4050085		
KEYWORDS				
SOURCE				
ORGANISM	Homo sapiens (human)			
REFERENCE	Homo sapiens Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eumetaria; Euarchontoglires; Primates; Catarrhini; Homnidae; Homo.			
AUTHORS	1 (bases 1 to 216387) Rowen,L., Madan,A., Qin,S., Abbasi,N., Dora,M., Ratcliffe,A., Madan,A., Dickhoff,R., Shaffer,T., James,R., Laaky,S. and Hood,L. Complete sequence of the gene for presentin 1 Unpublished 2 (bases 1 to 216387) Rowen,L. Direct Submission Submitted (30-NOV-1998) Department of Molecular Biotechnology, Box 357730 University of Washington, Seattle, Washington 98195, USA Sequencing methodology: high redundancy shotgun using plasmids. Interpersed Repeats were identified with RepeatMasker (available from http://ftp.genome.washington.edu/RM/RepeatMasker.html).			
JOURNAL	Location/Qualifiers			
AUTHORS	1..216387			
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JOURNAL	/mol_type="genomic DNA"			
COMMENT	/db_xref="taxon:9606"			
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	/clone_1lb="P. deJong RPC11"			
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	/rpt_family="AluYo"			
	750..1064			
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CDS	join(1465..1522,5979..6139,14890..15075,17363..17416, 17650..17733,21177..21463,23822..24044,24159..24248, 25789..25862,27305..27455,28794..29118,29491..29765, 30115..30213,33250..33297,37674..37766) /note="unknown, intron-exon boundaries defined by ESTs AI1476732, AA677603, AA677761, T40389, AA546306, AA147425, AA147333, AA910441, AA170655, and cDNA in L40392; the sequence in L40392 begins at 28812; the exons from 21177-21463 and 23822-24044 contain a repeat; only 1 EST, T40389 matches the exon from 21177-21463, since several 3' ESTs end in the beginning of this exon, and several 5' ESTs start at the end of this exon, it is possible that			
FEATURES	source			

	two genes are in this span; if so, the stop codon for the first gene could not be identified; the beginning and the end of this gene match a hypothetical "RNA recognition protein" from several species, based on blastx"
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repeat_region	/complement(3956..4121)
	/rpt_family="LMC4"
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	/rpt_family="AlusC"
repeat_region	/complement(4676..4975)
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repeat_region	/complement(5052..5163)
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Query Match      39.0%; Score 155, 8; DB 5; Length 216387;
Best Local Similarity 77.0%; Pred. No. 4, 2e-42;
Matches 218; Conservative 1; Mismatches 53; Indels 11; Gaps 2;

QY      81 ATGCGTGAATCCGAGCAGCTTCGGAGGCCAAGGTGGCCGAGTACCTGAGTCAAGAGA 140
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QY      201 TGGGCGATGGGGGACACACCGCTAGTCCGAGTACTAGCA-----GCCGAGATTGC 253
Db      195658 TGGGCGATGGGGGACACACCGCTAGTCCGAGTACTAGCA-----GCCGAGATTGC 195717
QY      254 AGTGAGCTGAGATGCGACAGTGAACCGAATACAGAT---CACAGATGAGCAGAGTG 309
Db      195718 CTTGACCTGAGAGGAGGAGGAGTGAACCAAGATTACTCAGCTTGTGATGACACAGCG 195777
QY      310 AGACKCGGTCTCAAAACACACAAACAAACAAACAAACCTA 352
Db      195778 AGACTCGGTCTCAAAACAAACAAACAAACAAACCTA 195820

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RESULT 37

AL732374/c
LOCUS
DEFINITION

Human DNA sequence from clone RP13-444K19 on chromosome X contains a mitochondrial ribosomal protein S18C (MRPS18C) pseudogene, the 3' end of the gene for a novel protein similar to PHD finger protein 2 PHF2 and a CpG island, complete sequence.

AL732374
VERSION
KEYWORDS
SOURCE
ORGANISM

AL732374.14 GI:23476649
HTG; CpG island; MRPS18C; PHF2.
Homo sapiens (human)
Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini; Homnidae; Homo.
1 (bases 1 to 224187)
Chapman, J.
Direct Submission

Submitted (13-MAY-2005) Wellcome Trust Sanger Institute, Hinxton, Cambridgeshire, CB10 1SA, UK. E-mail enquiries: vegas@sanger.ac.uk

Clone requests: clonerequests@sanger.ac.uk
On Oct 2, 2002 this sequence version replaced gi:23393869.
The following abbreviations are used to associate primary accession numbers given in the feature table with their source databases:

Emi, EMBL; SW, SWISSPROT; Tr, TREMBL; Wp, WORMPEP; Information on the WORMPEP database can be found at
http://www.sanger.ac.uk/Projects/C_elegans/wormpep This sequence was generated from part of bacterial clone contigs of human chromosome X, constructed by the Sanger Centre Chromosome X Mapping Group. Further information can be found at
http://www.sanger.ac.uk/HGP/ChrX

RP13-444K19 is from the library RP13-13.2 constructed by the group of Pieter de Jong. For further details see
http://www.chori.org/bacpac/home.htm
VECTOR: pBACe3.6

----- Genome Center
Center: Wellcome Trust Sanger Institute
Center code: SC
Web site: http://www.sanger.ac.uk
Contact: vegas@sanger.ac.uk

This sequence was finished as follows unless otherwise noted: all regions were either double-stranded or sequenced with an alternate chemistry or covered by high quality data (i.e., phred quality >= 30); an attempt was made to resolve all sequencing problems, such as compressions and repeats; all regions were covered by at least one subclone; and the assembly was confirmed by restriction digest.

FEATURES

except on the rare occasion of the clone being a YAC.

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CDS
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RPI-193M13₁s from the library RPCI-1 constructed by the group of
 Pieter de Jong. For further details see
<http://www.chori.org/bacpac/home.htm>
 VECTOR: pCYPAC2
 ----- Genome Center
 Center: Wellcome Trust Sanger Institute

```

/note="Clone left end: Rpt-193N13"
join(complement(5197..52970),complement(AL002272.1:1.19522..133076))
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 locus_tag="RP3_354N19.2-001"
 join(complement(52751..52970),complement(51672..52496),complement(10105..10201),complement(5197..5312),complement(AL002272.1:1.132996..133076),complement(AL002272.1:1.90573..90667),complement(AL002272.1:1.47072..47195),complement(AL002272.1:1.33567..43660),complement(AL002272.1:1.36090..36205),complement(AL002272.1:1.31977..32196),complement(AL002272.1:1.30718..30890),complement(AL002272.1:1.22625..22740),complement(AL002272.1:1.19522..22258))
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 join(complement(51672..52274),complement(10105..10201),complement(5197..5312),complement(AL002272.1:1.132996..133076),complement(AL002272.1:1.90573..90667),complement(AL002272.1:1.47072..47195),complement(AL002272.1:1.33567..43660),complement(AL002272.1:1.36090..36205),complement(AL002272.1:1.31977..32196),complement(AL002272.1:1.30718..30890),complement(AL002272.1:1.22625..22740),complement(AL002272.1:1.22132..22258))
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repeat_region 1981..2265
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repeat_region 14720..15036
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/rpt_family="Alu"
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repeat_region 16159..16198
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/rpt_family="MIR"

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Query Match 38.9%; Score 155.4; DB 5; Length 126462;
 Best Local Similarity 72.6%; Pred. No. 4.5e-42;
 Matches 217; Conservative 1; Mismatches 72; Indels 9; Gaps 1;

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Qy 81 ATGCGTAAATCCAGACCTCGGAGGCCAGGTGGCGGATCACTGAGTCAAGAGA 140
Db 93671 ACGCTTAATATCCAGACCTTTGGAAGGCCAAGTGGCGGATCACTGAGGCCAGGAT 93730
Qy 141 TCGAGACCATCTTGGCCAGACGTGTGAACCCCGCTTTACTTAAATACAAAATATAGC 200
Db 93731 TCGAGACCATCTTGGCCAGACGTGTGAACCCCGCTTTACTTAAATACAAAATATAGC 93790
Qy 201 TGGGATGCTGGGACACACCTGTGTCCAGCTCTCGAGAGCCGAGATTGCGTAGGC 260
Db 93791 CCGAGTGTGTGGGGGCGCTGTGTCCTCACTAGTGGAGGCGAGGTTGCGTAGGC 93850
Qy 261 TGAGATCGACAGAGTGAGCCGAATATCAGATACAGAGTGAGCGAGTGAACCCGCTCT 320
Db 93851 CGAGACCAAGCGCTTTGACATCTCCAGCTCGGGCGA-----CAGACGAGACTGTCT 93901
Qy 321 CAAAAACAACAACAAAAAACAATAGACATTTGTCATCTGCGGTTCCAG 379
Db 93902 CAACACAATTAACAAAAAACAAGTGAAGTCTTGCTTTGAGTTGGGGTCA 93960

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RESULT 40
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 LOCUS DEFINITION Homo sapiens chromosome 16 clone RP11-547D14, complete sequence.
 AC007616
 VERSION AC007616.5 GI:40018655
 KEYWORDS HTG.
 SOURCE Homo sapiens (human)
 ORGANISM Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini;

REFERENCE 1 (bases 1 to 132492)
 DOE Joint Genome Institute, Stanford Human Genome Center and Los
 Alamos National Laboratory.
 DIRECT SUBMISSION
 Unpublished

TITLE JOURNAL
 REFERENCE JOURNAL
 AUTHORS
 2 (bases 1 to 132492)
 Bruce, D., Mundt, M., Doggett, N., Munk, C., Saunders, E., Robinson, D.,
 Jones, M., Buckingham, J., Chasteen, L., Thompson, S., Goodwin, L.,
 Bryant, J., Tesmer, D., Melnick, J., Longmire, J., White, S., Tatam, O.,
 Campbell, C., Fawcett, J., Malbie, M., Bussod, M., Sutherland, R.,
 McMurtry, K., Han, C. and Deaven, L.

Direct Submission
 Submitted (20-MAY-1999) Center for Human Genome Studies, DOE Joint
 Genome Institute, Los Alamos National Laboratory, MS M888, Los
 Alamos, NM 87545, USA

REFERENCE 3 (bases 1 to 132492)
 AUTHORS DOE Joint Genome Institute.
 TITLE Direct Submission
 JOURNAL Submitted (06-JUL-2002) Production Sequencing Facility, DOE Joint Genome Institute, 2800 Mitchell Drive, Walnut Creek, CA 94598, USA
 REFERENCE 4 (bases 1 to 132492)
 AUTHORS Stanford Human Genome Center and Los Alamos National Laboratory.
 CONSRMT DOE Joint Genome Institute
 TITLE Direct Submission
 JOURNAL Submitted (18-DEC-2003) DOE Joint Genome Institute, 2800 Mitchell Drive, Walnut Creek, CA 94598, USA
 COMMENT On Dec 18, 2003 this sequence version replaced gi:21700556.
 Draft Sequence Produced by DOE Joint Genome Institute
 www.jgi.doe.gov
 Finishing Completed at Stanford Human Genome Center and Los Alamos National Laboratory
 www.hgsc.stanford.edu
 Quality: Phrap Quality >=40 100% of Sequence;
 Estimated Total Number of Errors is 0.

NOTE: This is not the entire sequence of the clone (entire sequence 168,9kb). It is clipped at the overlap with AC007613. The number of bases overlapped is 29296.

FEATURES
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 /organism="Homo sapiens"
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ORIGIN
 Query Match 38.9%; Score 155.4; DB 5; Length 132492;
 Best Local Similarity 73.1%; Pred. No. 4,6e-42;
 Matches 198; Conservative 1; Mismatches 72; Indels 0; Gaps 0;

83 GCCTTAATCCAGACTTCGGAGAGCCAGAGTGGCGGATCATCTGAGTCAAGATC 142
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 143 GAGACCATCTGGCCAGCATGTGTGAACCCGCTCTTAATAAATAACAAAAATAGCTG 202
 19680 GAGACCATCTGGCCAGCATGTGTGAACCCGCTCTTAATAAATAAATAAATAGCTG 19621
 203 GGCATGTGGGACACACTTGTAGTCCGACTTCTAGAGAGCCGAGATTGACGTGAGCTG 262
 19620 GGCCTGGTGGATGCGCTGTGCAATCCAGCTACTCAGAGAGGTGAGCGAGGAATTGCT 19561
 263 AGATGCGAGAGTGAAGCGAAATATCAGATCAAGAGTGAAGAGTGAAGAGTCA 322
 19560 TGAACCCAGAGAGAGGAGGAATTTGTTAAACCTTGGCGAGAGCGAGACTCCATCTCA 19501
 323 AAAACACACACAAAAACAAAAACATAA 353
 19500 AAAACACACACAAAAACAAAAACATAA 19470

RESULT 41
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 LOCUS Homo sapiens chromosome 11 clone RP11-753E7 map 11, WORKING DRAFT
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 AC021249.2 GI:7651924
 VERSION
 AC021249.2 GI:7651924
 KEYWORDS
 HTG; HTGS; PHASE1; HTGS_DRAFT.
 SOURCE
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 ORGANISM
 Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini; Hominoidea; Homo.
 REFERENCE
 1 (bases 1 to 145679)
 AUTHORS Birren, B., Linton, L., Nusbaum, C. and Lander, E.
 TITLE Homo sapiens chromosome 11, clone RP11-753E7

JOURNAL
 REFERENCE
 AUTHORS
 Unpublished
 2 (bases 1 to 145679)
 Birren, B., Linton, L., Nusbaum, C., Lander, E., Abraham, H., Allen, N., Anderson, S., Baldwin, J., Barna, N., Bastien, V., Beda, F., Boguslavsky, L., Bouhagalter, B., Brown, A., Burkett, G., Campopiano, A., Castle, A., Chappel, Y., Colangelo, M., Collins, S., Collymore, A., Cooke, P., Dekrellano, K., Dewar, K., Diaz, J. S., Dodge, S., Donno, M., Doyle, S., Ginde, S., Goyette, M., Graham, L., Galagan, J., Gardyna, S., Grant, G., Hagos, B., Heaford, A., Horton, L., Grand-Pierre, N., Iliev, I., Johnson, C., Jones, C., Kam, L., Karacas, A., Klein, J., Labocque, K., Lamazares, R., Landers, T., Lehoczy, J., Levine, R., Liu, C., Liu, G., Locke, K., MacDonald, P., Margis, N., McCarthy, M., McKean, P., McGuck, A., McKernan, K., McPheters, R., Melchior, J., Meneus, L., Minova, T., Miranda, C., Mlenka, V., Morrow, J., Murphy, T., Naylor, J., Norman, C. H., O'Connor, T., O'Donnell, P., O'Neill, D., Oliver, T. M., Oliver, J., Peterson, K., Pierre, N., Plesani, C., Pollara, V., Raymond, C., Riley, R., Rogov, P., Rothman, D., Roy, A., Santos, R., Schauer, S., Severy, P., Spencer, B., Stange-Thomann, N., Stojanovic, N., Sudhanian, A., Talamas, J., Tealaye, S., Theodore, J., Tittell, A., Travers, M., Triggillo, J., Vassiliev, H., Viel, R., Vo, A., Wilson, B., Wu, X., Wyman, D., Ye, W. J., Young, G., Zainoun, J., Zimmer, A. and Zody, M.

Direct Submission
 Submitted (28-MAR-2000) Whitehead Institute/MIT Center for Genome Research, 320 Charles Street, Cambridge, MA 02141, USA
 On Apr 27, 2000 this sequence version replaced gi:7331619.
 All repeats were identified using RepeatMasker:
 Smit, A.F.A. & Green, P. (1996-1997)
 http://ftp.genome.washington.edu/RM/RepeatMasker.html

Genome Center
 Center: Whitehead Institute/ MIT Center for Genome Research
 Center code: WIR
 Web site: http://www-seq.wi.mit.edu
 Contact: sequence.submissions@genome.wi.mit.edu

Project Information
 Center project name: 18860
 Center clone name: 753_E7

Sequencing Vector: M13; M77815; 100% of reads
 Chemistry: Dye-terminator Big Dye; 100% of reads
 Assembly program: Phrap; version 0.960731
 Consensus quality: 129416 bases at least Q40
 Consensus quality: 137400 bases at least Q30
 Consensus quality: 140385 bases at least Q20
 Insert size: 154000; agarose-fp
 Insert size: 142279; sum-of-coverage
 Quality coverage: 3.9 in Q20 bases; agarose-fp
 Quality coverage: 4.2 in Q20 bases; sum-of-coverage

NOTE: This is a 'working draft' sequence. It currently consists of 35 contigs. The true order of the pieces is not known and their order in this sequence record is arbitrary. Gaps between the contigs are represented as runs of N, but the exact sizes of the gaps are unknown. This record will be updated with the finished sequence as soon as it is available and the accession number will be preserved.

1 1125: contig of 1125 bp in length
 * 1126 1225: gap of 100 bp
 * 1126 2425: contig of 1201 bp in length
 * 2427 2526: gap of 100 bp
 * 2527 3864: contig of 1338 bp in length
 * 3865 3964: gap of 100 bp
 * 3965 5223: contig of 1259 bp in length
 * 5224 5323: gap of 100 bp
 * 5324 7019: contig of 1696 bp in length
 * 7020 7119: gap of 100 bp
 * 7120 9177: contig of 2058 bp in length
 * 9178 9277: gap of 100 bp
 * 9278 11580: contig of 2303 bp in length
 * 11581 11680: gap of 100 bp
 * 11681 13580: contig of 1900 bp in length

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* 13581 13680: gap of 100 bp
* 13681 15409: contig of 2729 bp in length
* 16410 16509: gap of 100 bp
* 16510 19008: contig of 2499 bp in length
* 19009 19108: gap of 100 bp
* 19109 21633: contig of 2525 bp in length
* 21634 21733: gap of 100 bp
* 21734 24035: contig of 2302 bp in length
* 24036 24135: gap of 100 bp
* 24136 26117: contig of 1982 bp in length
* 26118 28765: contig of 2548 bp in length
* 28766 28865: gap of 100 bp
* 28866 31673: contig of 2808 bp in length
* 31674 34537: contig of 2764 bp in length
* 34538 37137: contig of 2500 bp in length
* 37138 37237: gap of 100 bp
* 37238 40767: contig of 3530 bp in length
* 40768 44314: contig of 3447 bp in length
* 44315 46751: contig of 2337 bp in length
* 46752 49744: contig of 2893 bp in length
* 49745 53918: contig of 4074 bp in length
* 53919 57127: contig of 3109 bp in length
* 57128 61720: contig of 4493 bp in length
* 61721 66820: gap of 100 bp
* 66821 66853: contig of 4833 bp in length
* 66854 70529: contig of 3776 bp in length
* 70530 75419: contig of 4790 bp in length
* 75420 81973: gap of 6454 bp in length
* 81974 82074 91465: contig of 9392 bp in length
* 91466 91565: gap of 100 bp
* 91566 99702: contig of 8137 bp in length
* 99703 99802: gap of 100 bp
* 99803 107085: contig of 7283 bp in length
* 107086 107185: gap of 100 bp
* 107186 114948: contig of 7763 bp in length
* 114949 115048: gap of 100 bp
* 115049 123618: contig of 8570 bp in length
* 123619 131968: contig of 8250 bp in length
* 131969 132069: gap of 100 bp
* 132069 145679: contig of 13611 bp in length.
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gap 5324. .7019
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Matches 203; Conservative 1; Mismatches 62; Indels 1; Gaps 1;

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DB 8464 TCGAGATCAAGCTGAGCCCAATGTTGAACCCCGCTTTACTAAAAATCAAAAAATAGC 8405
QY 201 TGGGATGTTGGCAACACCTGTAGTCCAGCTACTCAGAGCCGAGATTGCAATGAC 260
DB 8404 TGGGATGTTGGCCCAAGCTGTAGTCCAGCTACTCAGAGCCGAGATTGCAATGAC 8345
QY 261 TGAATCGCAGAGTGAAGCCCAATCAGATCAGAGTGAAG-CAGGTGAGACCGGTC 319
DB 8344 CTGTAACCCAGAGCGGAGGCTCAGATGAGCCCAAGATCGGACAGCAAGATTCCGTC 8285
QY 320 TCAAAACCAACACAAAAAACA AAAA 346
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RESULT 42
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DEFINITION
ACCESSION AP001024 GI:3240066
VERSION AP001024.6
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KEYWORDS HTG.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini; Homnidae; Homo.

REFERENCE
AUTHORS Hattori, M., Ishii, K., Toyoda, A., Taylor, T. D., Hong-Seog, P., Fujiyama, A., Yada, T., Totoki, Y., Matsunabe, H. and Sakaki, Y.
TITLE Homo sapiens genomic DNA
JOURNAL Published Only in Databases (2000)
AUTHORS 2 (bases 1 to 183444)
Hattori, M., Ishii, K., Toyoda, A., Taylor, T. D., Hong-Seog, P., Fujiyama, A., Yada, T., Totoki, Y., Matsunabe, H. and Sakaki, Y.
JOURNAL Direct Submission
Submitted (05-JAN-2000) Mashima Hattori, The Institute of Physical and Chemical Research (RIKEN), Genomic Sciences Center (GSC); 1-7-22 Suenhiro-chou, Tsurumi-ku, Yokohama, Kanagawa 230-0045, Japan (E-mail:hattori@gs.c.riken.go.jp, URL:http://hgp.gs.c.riken.go.jp/, Tel:81-45-503-9111, Fax:81-45-503-9170)
COMMENT On Jul 1, 2003 this sequence version replaced gi:31790714.

FEATURES
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/organism="Homo sapiens"
/mol_type="genomic DNA"
/db_xref="taxon:9606"
/chromosome="11"
/map="11q"
/clone="RP11-832N8"

ORIGIN

Query Match 38.9% Score 155.4; DB 5; Length 183444;
Best Local Similarity 76.0%; Pred. No. 5,4e-42;
Matches 203; Conservative 1; Mismatches 62; Indels 1; Gaps 1;

QY 81 ATGCTGTATATCCAGACCTTCGGAGGCGCAAGTGGCGGATCACTGAGGTCAAGAGA 140
DB 173708 ATGCTGTATATCCAGACCTTCGGAGGCGCAAGTGGCGGATCACTGAGGTCAAGAGA 173767
QY 141 TCGAAGACCTCTCTGGCCCAAGTGTGAAGAACCCGCTTTTCAAAAAATCAAAAAATAGC 200
DB 173768 TCGAAGACCTCTCTGGCCCAAGTGTGAAGAACCCGCTTTTCAAAAAATCAAAAAATAGC 173827
QY 201 TGGGATGATGCGACACACCTGTAGTCCAGCTACTGACGAGCGGAGATTCCAGTAGAC 260
DB 173828 TGGGATGATGCGACACACCTGTAGTCCAGCTACTGACGAGCGGAGATTCCAGTAGAC 173887
QY 261 TGAGATCGCAGAGTGGCGCAATCAAGATCAAGAGTAGAG-CAGAGTGAAGACKCGTC 319
DB 173888 CTGGAACCCAGAGGCGGAGGCTGCAAGTGAAGTCCGACAGACAGATTCCTGTC 173947
QY 320 TCAAAAAACACCAAAAAACAAAAA 346
DB 173948 TCAAAAAACACCAAAAAACAAAAAAGAGAA 173974

RSULT 43
AC018423/c 202138 bp DNA linear HTG 06-MAY-2001
LOCUS Homo sapiens chromosome 11 clone RP11-531B6 map 11, WORKING DRAFT
DEFINITION
SEQUENCE, 22 unordered pieces.
AC018423
AC018423 4 GI:10045404
KEYWORDS HTG; HTGS_PHASE1; HTGS_DRAFT.
SOURCE
ORGANISM Homo sapiens (human)
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini; Homnidae; Homo
REFERENCE
AUTHORS Birren, B., Linton, L., Nusbaum, C. and Lander, E.
TITLE Homo sapiens chromosome 11, clone RP11-531B6
JOURNAL Unpublished

REFERENCE
AUTHORS
2 (bases 1 to 202138)
Birren, B., Linton, L., Nusbaum, C., Lander, E., Abraham, H., Allen, N., Anderson, S., Baldwin, J., Barna, N., Beckerly, R., Beda, F., Boguslavsky, L., Boukhgalter, B., Brown, A., Castle, A., Colangelo, M., Collins, S., Collymore, A., Cooke, P., Dearlano, K., Dewar, K., Domino, M., Doyle, M., Feneiro, J., Ferreira, P., Fitzhugh, W., Forrest, C., Gage, D., Galagan, J., Gardyna, S., Grant, G., Hagos, B., Heath, A., Horton, L., Howland, J., C., Johnson, R., Jones, C., Kann, L., Kartas, A., Klein, J., Lander, T., Lehotzky, J., Lien, C., Locke, K., Macdonald, P., Margulis, N., McEwan, P., McGuirk, A., McKernan, K., Meltrina, J., Morrow, J., Naylor, J., Norman, C. H., O'Connor, T., O'Donnell, P., Peterson, K., Pierre, N., Pollara, V., Riley, R., Rothman, D., Roy, A., Santos, R., Severy, P., Strange-Thomann, N., Stojanovic, N., Sudramanian, A., Talamas, J., Testaye, S., Theodore, J., Tirrell, A., Vassiliev, H., Viet, R., Vo, A., Wu, X., Wyman, D., Ye, W. J., Zimmer, A. and Zody, W.

JOURNAL Direct Submission
Submitted (10-DEC-1999) Whitehead Institute/MIT Center for Genome Research, 320 Charles Street, Cambridge, MA 02141, USA
On Sep 9, 2000 this sequence version replaced gi:6649385.
All repeats were identified using RepeatMasker:
Smit, A.F.A. & Green, P. (1996-1997)
http://ftp.genome.washington.edu/RM/RepeatMasker.html

COMMENT
Center: Whitehead Institute/ MIT Center for Genome Research
Center code: WIBR
Web site: http://www.seq.wi.mit.edu
Contact: sequence_submissions@genome.wi.mit.edu
Project Information
Center project name: L3744
Center clone name: 531.E.6
Summary Statistics
Sequencing vector: M13; M7815; 100% of reads
Chemistry: Dye-terminator Big Dye; 100% of reads
Assembly program: Phrap; version 0.960731
Consensus quality: 186769 bases at least Q40
Consensus quality: 198059 bases at least Q20
Insert size: 21000; agarose-fp
Insert size: 200038; sum-of-contigs
Quality coverage: 4.4 in Q20 bases; agarose-fp
Quality coverage: 4.6 in Q20 bases; sum-of-contigs

NOTE: This is a 'working draft' sequence. It currently consists of 22 contigs. The true order of the pieces is not known and their order in this sequence record is arbitrary. Gaps between the contigs are represented as runs of N, but the exact sizes of the gaps are unknown. This record will be updated with the finished sequence as soon as it is available and the accession number will be preserved.

1
2851: contig of 2851 bp in length
2852 2951: gap of 100 bp
2952 4977: contig of 2026 bp in length
4978 5077: gap of 100 bp
5078 8300: contig of 3223 bp in length
8301 8400: gap of 100 bp
8401 10751: contig of 2351 bp in length
10752 10851: gap of 100 bp
10852 14527: contig of 3676 bp in length
14528 14627: gap of 100 bp
14628 53554: contig of 38927 bp in length
53555 53555: gap of 100 bp
53556 59471: contig of 5817 bp in length
59472 59571: gap of 100 bp
59572 65624: contig of 6053 bp in length
65625 65724: gap of 100 bp
65725 72213: contig of 6489 bp in length
72214 72313: gap of 100 bp
72314 76349: contig of 4036 bp in length
76350 76450: gap of 100 bp
76451 82044: contig of 5595 bp in length
82045 82144: gap of 100 bp

FEATURES	source
*	83145: contig of 8122 bp in length
*	90317: gap of 100 bp
*	96732: contig of 6316 bp in length
*	96733: gap of 100 bp
*	96833: contig of 9755 bp in length
*	106558: gap of 100 bp
*	106558: contig of 7758 bp in length
*	114416: gap of 100 bp
*	114515: contig of 8229 bp in length
*	122745: gap of 100 bp
*	122844: contig of 9330 bp in length
*	132174: gap of 100 bp
*	132274: contig of 12216 bp in length
*	133175: gap of 100 bp
*	133275: contig of 10084 bp in length
*	144590: gap of 100 bp
*	144591: contig of 10084 bp in length
*	145574: gap of 100 bp
*	155675: contig of 18202 bp in length
*	173877: gap of 100 bp
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|vector_side:left"
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|estimated_length=100
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8401..10751
|note="assembly_fragment"
|one_end:SP6
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10852..14527
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14528..14627
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gap	114416. .114515 /estimated_length=100 114516. .122744 /note="assembly_fragment"
misc_feature	122745. .122844 /estimated_length=100 122845. .132174 /note="assembly_fragment"
gap	132175. .132274 /estimated_length=100 132275. .144490 /note="assembly_fragment"
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Best Local Similarity	76.0%	Pred. No. 5.6e-42;		
Matches 203; Conservative	1;	Mismatches 62;	Indels 1;	Gaps 1;

OY	81	ATGCTCTGTAATCCCGACACTTCGGAGGCGCAAGGTGGCGGATTCACCTGAGGTCAAGAGA	140
Db	102362	ATGCTCTGTAATCCCGACACTTCGGAGGCGCAAGGCGGAGTCACTTGAAGTCAGGAAGT	102303
OY	141	TCGAGACCATCCTGGCCCAACATGTGTGAACCCCGCTCTTTACTTAAATAACAAAAATAGC	200
Db	102302	TCGAGATCAGTCTACCAACATGTGTGAACCTTGTCTCTACTTAAAAATCAAAAGATTAGC	102243
OY	201	TGGCATGTGTGGCACACACTGTACTGCCAGTACTTCAGAGACCGGAGTTTGCAGTGAGC	260
Db	102242	TGGGTGTGTGGCCCAACGCTGTGTATCCAGTACTTCAGAGACGTGAGACAGGAGATGC	102183
OY	261	TGAGATCCAGAGTGTAGCCGGAATACAGATACAGAGTGAG-CAGAGTGAGACKCCGTC	319
Db	102182	CTTGAAACCCAGGAGGCGGAGGCTGCAAGTGAGCCAAAGTGGCAGACGAAGATTCCGTC	102123
OY	320	TCAAAAACACACAAAAACAAAAA	346
Db	102122	TCAAAAACAAAAACAAAAACGAGAA	102096

RESULT	44
AC009489	
LOCUS	204493 bp DNA linear PRI 04-FEB-2003
DEFINITION	Homo sapiens chromosome 16 clone CTF-3808G3, complete sequence.
ACCESSION	AC009489
VERSION	AC009489.2 GI:28201477
KEYWORDS	HTG.
SOURCE	Homo sapiens (human)

ORGANISM Homo sapiens
Eukaryota; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini;
Homidae; Homo.

REFERENCE 1 (bases 1 to 204493)
AUTHORS DOE Joint Genome Institute, Stanford Human Genome Center and Los
Alamos National Laboratory.
TITLE Direct Submission
JOURNAL Unpublished
REFERENCE 2 (bases 1 to 204493)
AUTHORS DOE Joint Genome Institute.
TITLE Direct Submission
JOURNAL Submitted (15-NOV-2001) Production Sequencing Facility, DOE Joint
Genome Institute, 2800 Mitchell Drive, Walnut Creek, CA 94598, USA
3 (bases 1 to 204493)
AUTHORS DOE Joint Genome Institute, Stanford Human Genome Center and Los
Alamos National Laboratory.
TITLE Direct Submission
JOURNAL Submitted (04-FEB-2003) DOE Joint Genome Institute, 2800 Mitchell
Drive, Walnut Creek, CA 94598, USA
On Feb 4, 2003 this sequence version replaced gi:16930905.
Draft Sequence Produced by DOE Joint Genome Institute
www.jgi.doe.gov
Finishing Completed at Stanford Human Genome Center and Los Alamos
National Laboratory
www.hgsc.stanford.edu
Quality: Phrap Quality >=40 99.8% of Sequence;
Estimated Total Number of Errors is 0.4.
NOTE: Shatter libraries failed to verify the dinucleotide repeat
region 65514-65789. Unsure number of repeat copies 65514-65789.
Forced join at 65675.

FEATURES
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/clone="CTD-308863"
65514..65789
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repeat copies 65514-65789. Forced join at 65675."

ORIGIN
Query Match 38.9%; Score 155.4; DB 5; Length 204493;
Best Local Similarity 73.1%; Pred. No. 5.7e-42;
Matches 198; Conservative 1; Mismatches 72; Indels 0; Gaps 0;

83 GCGGTATATCCAGACTTCGGAGGCCAAGGTGGCGGATCACTGAGGTCAAGAGATC 142
|||||
14479 GCGGTATATCCAGACTTCGGAGGCCAAGGTGGCGGATCACTGAGGTCAAGAGATC 14538
|||||
143 GAGACCATCTGGCCCAATGCTGAACCCCTCTTACTTAAATAACAAAATAGCTG 202
14539 GAGACCATCTGGCCCAATGCTGAACCCCTCTTACTTAAATAACAAAATAGCTG 14558
203 GCGGTATATCCAGACTTCGGAGGCCAAGGTGGCGGATCACTGAGGTCAAGAGATC 262
14599 GCGGTATATCCAGACTTCGGAGGCCAAGGTGGCGGATCACTGAGGTCAAGAGATC 14658
263 AGATGGAGAGTGAAGCCGAATACAGATCAAGAGTGAAGAGTGAAGAGTGAAGAGTCA 322
14659 TGAATCCAGAGAGAGAGAGATGCTTGAACCTCTGGCGGAGAGAGAGTCACTCA 14718
323 AAAAC 353
14719 AAAAC 14749

RESULT 45
LOCUS AL355388 205463 bp DNA linear PRI 18-MAY-2005
DEFINITION Human DNA sequence from clone RP11-336K24 on chromosome 1 Contains

the 5' end of the RIT1 gene for Ras-like without CAAX 1, the gene
for a novel protein (KIAA0907), the ARHGAP2 gene for Rho/rac
guanine nucleotide exchange factor (GEF) 2, four novel genes, the
SSR2 gene for signal sequence receptor beta (translucan-associated
protein beta), the Clorf6 gene for chromosome 1 open reading frame
6, the gene for mitogen-activated protein-binding
protein-interacting protein (MAPBIP), the RAB25 gene for RAB25
(member RAS oncogene family), the 5' end of the LMNA gene for Lamin
A/C and three CpG islands, complete sequence.

ACCESSION AL355388
VERSION AL355388.30, GI:29367464
KEYWORDS HTG; ARHGAP2; Clorf6; KIAA0907; LMNA; MAPBIP; RAB25; RIT1; SSR2.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini;
Homidae; Homo.

REFERENCE 1 (bases 1 to 205463)
AUTHORS Hall, R.
TITLE Direct Submission
JOURNAL Submitted (13-MAY-2005) Wellcome Trust Sanger Institute, Hinxton,
Cambridgeshire, CB10 1SA, UK. E-mail enquiries: vegas@sanger.ac.uk
Clone requests: clonerequest@sanger.ac.uk
On Mar 29, 2003 this sequence version replaced gi:28446036.
The following abbreviations are used to associate primary accession
numbers given in the feature table with their source databases:
Emi., EMBL; Sw., SWISSPROT; Tr., TREMBL; Wp., WORMPEP; Information
on the WORMPEP database can be found at
http://www.sanger.ac.uk/Projects/C_elegans/wormpep This sequence
was generated from part of bacterial clone contigs of human
chromosome 1, constructed by the Sanger Centre Chromosome 1 Mapping
Group. Further information can be found at
http://www.sanger.ac.uk/HGP/Chr1
Of Pieter de Jong. For further details see
http://www.choi.org/bacpac/home.htm
VECTOR: pBAC3.6

----- Genome Center
Center: Wellcome Trust Sanger Institute
Center code: SC
Web site: http://www.sanger.ac.uk
Contact: vegas@sanger.ac.uk

This sequence was finished as follows unless otherwise noted: all
regions were either double-stranded or sequenced with an alternate
chemistry or covered by high quality data (i.e., phred quality >=
30); an attempt was made to resolve all sequencing problems, such
as compressions and repeats; all regions were covered by at least
one subclone; and the assembly was confirmed by restriction digest,
except on the rare occasion of the clone being a YAC.

FEATURES
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1
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complement(4130..4186),
complement(AL139128..24:140824..141015),
complement(AL139128..24:134321..137131))

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complement(AL139128.24:140824..141015),
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match: CDNAS: Em:AF084462.1 Em:U71203.1 Em:U78165.1
Em:Y07566.1"
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complement(AL139128.24:141116..141123),
complement(AL139128.24:140824..141015),
complement(AL139128.24:136911..137131))
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/locus_tag="RP11-10106.4-003"
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complement(AL139128.24:141244..141317),
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complement(AL139128.24:140824..141015),
complement(AL139128.24:136911..137131))
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/product="Ras-like without CAAX 1"
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complement(AL139128.24:140824..141015),
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/locus_tag="RP11-10106.4-002"
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complement(AL139128.24:141244..141317),
complement(AL139128.24:140824..141015),
complement(AL139128.24:136787..137131))
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/join(complement(4558..4565),complement(4336..4484),
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complement(AL139128.24:141244..141317),
complement(AL139128.24:140824..141015),
complement(AL139128.24:136787..137131))
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/product="Ras-like without CAAX 1"
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complement(AL139128.24:140824..141015),
complement(AL139128.24:136901..137131))
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/product="Ras-like without CAAX 1"
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/db_xref="GOA:Q5VY89"
/db_xref="InterPro:IPR001806"
/db_xref="InterPro:IPR002041"
/db_xref="InterPro:IPR003577"
/db_xref="InterPro:IPR003578"
/db_xref="InterPro:IPR003579"
/db_xref="InterPro:IPR005225"
/db_xref="UniProtKB/TrEMBL:O5VY89"
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GAGYVKSAMTQPIFSRPPEDHPDITDAKIRIRIDDEPANLIDLTAGAEFTTAM
RQYVRABGFLICYSIDRDSFHVREKQOLIYRVRTDTPVYLVNKSJDKQIOL
VTEKGLALAREFSCPFETSAARYYIDVDFHALVREIRKEKAVLAIEKSKSPKNS
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/join(complement(4336..4441),complement(4130..4186),

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complement(AL139128.24:140824..141015),
complement(AL139128.24:136901..137131))
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/product="Ras-like without CAAX 1"
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/db_xref="InterPro:IPR005225"
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complement(join(6723..8000,9553..9644,10305..10417,
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23392..23520,27313..27529,28016..28077))
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Em:BC006621.1 Em:BC062637.1"
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/locus_tag="RP11-336K24.1-001"
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15055..15308,15541..15598,17307..17367,19312..19523,
20356..20519,20784..20836,22976..23055,23392..23520,
27313..27529,28016..28079))
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/locus_tag="RP11-336K24.1-003"
/complement(join(6725..9644,10305..10417,11179..11352,
15055..15308,15541..15598,17307..17367,19312..19523,
20356..20519,20784..20836,22976..23055,23392..23520,
27313..27529,28016..28079))
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/locus_tag="RP11-336K24.1-003"
/product="novel protein"
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polyA_site
Query Match 38.9%; Score 155.4; DB 5; Length 205463;
Best Local Similarity 74.7%; Pred. No. 5.76-42;
Matches 207; Conservative 1; Mismatches 67; Indels 2; Gaps 1;

polyA_site
81 ATGCTGTAATCCGACGACCTTCGGAGGCGCAAGTGGCGGATACCTGAGTCAAGGA 140
177219 ACGGCTGTAATCCGACGACCTTCGGAGGCGCAAGTGGCGGATACCTGAGTCAAGGA 177278
141 TCGAGACCATCTCTGCGCAACATGTTGAACCCCGTCTTTACTAAATAACAAAATAGC 200
177279 TCGAGACCATCTCTGCGCAACATGTTGAACCCCGTCTTTACTAAATAACAAAATAGC 177338
201 TGGGATGATGGACACACCTGTAGTCCAGCTACTCAGAGCCGAGATTGCAGTGAGC 260
177339 CGAGCATGTGTGCGAGACCGCTGTAGTCCAGCTACTCAGAGCGGTGAGACAG--GAGAT 177396
261 TGAATGCGCAAGTGAAGCGGAATTCACAGATCACAGAGTGAGAGAGTGAAGCCGGCT 320
177397 TGCTTGAACCCGAGGAGGGGAAGTTGCGATGATGATGCGAGAGTGAAGTCCGCTCC 177456

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DB	177457	CCCCCAAAAAAAAAACAAAAACAAAAACAACAAACAAA	177493
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DEFINITION	Human sapiens chromosome 16 clone CTD-2169M19, complete sequence.		
ACCESSION	AC026423		
VERSION	AC026423.9		
KEYWORDS	HTG.		
SOURCE	Human sapiens (human)		
ORGANISM	Human sapiens (human)		
REFERENCE	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini; Homiidae; Homo.		
AUTHORS	1 (bases 1 to 116130)		
TITLE	DOE Joint Genome Institute, Stanford Human Genome Center and Los Alamos National Laboratory.		
JOURNAL	Direct Submission		
REFERENCE	2 (bases 1 to 116130)		
AUTHORS	DOE Joint Genome Institute.		
TITLE	Direct Submission		
JOURNAL	Submitted (22-MAR-2000) Production Sequencing Facility, DOE Joint Genome Institute, 2800 Mitchell Drive, Walnut Creek, CA 94598, USA		
REFERENCE	3 (bases 1 to 116130)		
AUTHORS	DOE Joint Genome Institute.		
TITLE	Direct Submission		
JOURNAL	Submitted (04-JUN-2002) Production Sequencing Facility, DOE Joint Genome Institute, 2800 Mitchell Drive, Walnut Creek, CA 94598, USA		
REFERENCE	4 (bases 1 to 116130)		
AUTHORS	DOE Joint Genome Institute, Stanford Human Genome Center and Los Alamos National Laboratory.		
TITLE	Direct Submission		
JOURNAL	Submitted (18-MAR-2003) DOE Joint Genome Institute, 2800 Mitchell Drive, Walnut Creek, CA 94598, USA		
COMMENT	On Mar 18, 2003 this sequence version replaced gi:18057078. Direct Sequence Produced by DOE Joint Genome Institute www.jgi.doe.gov		
FEATURES	Finishing Completed at Stanford Human Genome Center and Los Alamos National Laboratory www.sngc.stanford.edu Quality: Phrap Quality >=40 100% of Sequence; Estimated Total Number of Errors is 0.		
SOURCE	Location/Qualifiers 1. 116130 /organism="Homo sapiens" /mol_type="genomic DNA" /db_xref="taxon:9606" /chromosome="16" /clone="CTD-2169M19"		
ORIGIN			
Query Match	38.8%; Score 155.2; DB 5; Length 116130;		
Best Local Similarity	74.5%; Pred. No. 5.1e-42;		
Matches 210; Conservative 1; Mismatches 64; Indels 7; Gaps 1;			
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DB	141	TCGAGACCATCTGGCCAAACATGTGGTAACCCCGCTTTAATAAAATACAAAAATAGC	200
DB	11367	TCGAGACCAAGCTGGCCAAACATGTGAGACCTCATCTTACTTAATAAAATACAAAAATTTAGC	1142
DB	201	TGGGATGATGGGACACACACTGTGATGCCAGACTACTCAGAGCCGGAATTCAGTAGC	260
DB	11427	CGGGCTTGATGGAGATGTGCTCTGTAATCCCACTACTCAGAGGCTGAGGACGAGAGAGT	1148
DB	261	TGAATGCGCAAGTGAAGCCGAATTCACAGAT-----CAGAGTAGCAGAGTAGAC	313

[illegible]

TITLE	Direct Submission
JOURNAL	Submitted (30-SEP-2001) Human Genome Sequencing Center, Department of Molecular and Human Genetics, Baylor College of Medicine, One Baylor Plaza, Houston, TX 77030, USA
REFERENCE	4 (bases 1 to 177640)
AUTHORS	Morley,K.C.
TITLE	Direct Submission
JOURNAL	Submitted (02-OCT-2001) Human Genome Sequencing Center, Department of Molecular and Human Genetics, Baylor College of Medicine, One Baylor Plaza, Houston, TX 77030, USA
REFERENCE	5 (bases 1 to 177640)
AUTHORS	Morley,K.C.
TITLE	Direct Submission
JOURNAL	Submitted (07-MAR-2002) Human Genome Sequencing Center, Department of Molecular and Human Genetics, Baylor College of Medicine, One Baylor Plaza, Houston, TX 77030, USA
REFERENCE	6 (bases 1 to 177640)
AUTHORS	Morley,K.C.
TITLE	Direct Submission
JOURNAL	Submitted (26-JUN-2002) Human Genome Sequencing Center, Department of Molecular and Human Genetics, Baylor College of Medicine, One Baylor Plaza, Houston, TX 77030, USA
REFERENCE	7 (bases 1 to 177640)
AUTHORS	Morley,K.C.
TITLE	Direct Submission
JOURNAL	Submitted (23-AUG-2002) Human Genome Sequencing Center, Department of Molecular and Human Genetics, Baylor College of Medicine, One Baylor Plaza, Houston, TX 77030, USA
REFERENCE	8 (bases 1 to 177640)
AUTHORS	Morley,K.C.
TITLE	Direct Submission
JOURNAL	Submitted (22-OCT-2002) Human Genome Sequencing Center, Department of Molecular and Human Genetics, Baylor College of Medicine, One Baylor Plaza, Houston, TX 77030, USA
COMMENT	On Oct 22, 2002 this sequence version replaced gi:15809129. INFORMATION: http://www.hgsc.bcm.tmc.edu/ or email gc-help@bcm.tmc.edu

CLONE LENGTH: This sequence does not necessarily represent the entire insert of this clone. Overlapping regions of clones are only sequenced and submitted once, so the sequence for the remainder of the insert may be found in the record for the adjacent clones. Overlapping clones are noted at the beginning and end of the Features listing.

ANNOTATION OF FEATURES:

STSs are identified using ePCR (Genome Res. 7:541-550) searches of a local database that includes entries from dbSTS, GDB, and local mapping efforts.

Repeats are identified using RepeatMasker (A. Smit and P. Green, unpublished.) for Human and Mouse sequences.

Genes and Region of sequence similarity are identified by BLAST (Nuc. Acids Res. 25:3389-3402) similarity (expect < 1e-34) to the EST and cDNA sequences. Genes demonstrate at least two exons flanked by consensus splice sites that maintained sequence continuity across the splice junctions. Sequences that are not identical matches are annotated as similar.

SEQUENCING READ COVERAGE:Sequencing is completed to a minimum standard of double strand coverage with a minimum of 2 clones and 2 reads with no ambiguities or 2 chemistries with a minimum of 2 clones and 3 reads with no ambiguities. If the sequence quality for a region does not meet this standard, it will be indicated in the annotation as Low Coverage.

QUALITY OF INDIVIDUAL BASES:This sequence meets stringent quality standards - estimated error rate less than 1 per 10,000 bases. Reports of lowest quality individual bases and measures of base quality are listed below. Description of the metrics can be found at URL:
<http://www.hgsc.bcm.tmc.edu:8088/quality.info/genbank.annotation.html>.

Location/Qualifiers

Query	DB	141	TCGACACCCTCTGGCCACATGCTGAACCCCGCTTTACTTAAATATCAAAAAATAC	200
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Matches 196; Conservative 1;	Mismatches 69; Indels 0; Gaps 0;			
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DB	97632	ATGCGCTTAATCCAGCACTTCGGAGGCCAGGAAGAGTGGATCACTGAGTCAAGAGT	97573	
141	TCGACACCCTCTGGCCACATGCTGAACCCCGCTTTACTTAAATATCAAAAAATAC	200		

Db	97512	TCAAGACAGCCTGGCCAAACATGTTGAACCTTGTCTCTAATTAACTACAAAATTAC	97513
Ox	201	TGGCATGTGTGGCACACACCTGTATGTCCACGACTACTCAGAGCCGAGATTGCATGTAC	260
Db	97512	CGGCTCTGTGTGACTCACACCTGTGTGTGCTCCAGCTACTCAGAGGCTGACGAGAAATG	97453
Ox	261	TGAATATGGCAGAGTACGAGCCGAAATACAGATATCAGAGTACGAGACTGAGACCCGTCT	320
Db	97452	CTTGAATCCAGAGAGCCGAGAGTGTACAGAGCAGACTGTGGTGTATGAGCAATTA	97393
Ox	321	CAAAACACACACAAAACAAAACAAA 346	
Db	97392	CAAAACACAAAACAAAACAAAACAAA 97367	
RESULT 48			
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LOCUS			
DEFINITION	AC068929	178079 bp	DNA linear HTG 07-DEC-2000
			Homo sapiens chromosome 1 clone RP11-206C14 map 1, WORKING DRAFT
			SEQUENCE, 11 unordered pieces.
AC068929			
AC068929.3			GI:11597096
HTG			HTG PHASE1: HTGS_DRAFT.
KEYWORDS			Homo sapiens (human)
SOURCE			Homo sapiens
ORGANISM			Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini; Homnidae; Homo.
REFERENCE			1 (bases 1 to 178079)
AUTHORS			Birren, B., Linton, L., Nusbaum, C. and Lander, E.
TITLE			Homo sapiens chromosome 1, clone RP11-206C14
JOURNAL			Unpublished
REFERENCE			2 (bases 1 to 178079)
AUTHORS			Birren, B., Linton, L., Nusbaum, C., Lander, E., Abraham, H., Allen, N., Anderson, S., Baldwin, J., Barna, N., Baerlen, V., Beda, F., Boguslavsky, L., Boungalter, B., Brown, A., Burkett, G., Campoliano, A., Castle, A., Choepel, Y., Colangelo, M., Collins, S., Collamore, A., Cooke, P., Deatrelino, K., Dewar, K., Diaz, J.S., Dodge, S., Domino, M., Doyle, M., Ferreira, P., Fitzhugh, W., Gage, D., Galagan, J., Gardyna, S., Glend, S., Goyette, M., Graham, L., Grand-Pierre, N., Grant, G., Hagos, B., Heatford, A., Horton, L., Howland, J.C., Iliev, I., Johnson, R., Jones, C., Kann, L., Karatas, A., Klein, J., Labocque, K., Lamazares, R., Landers, T., Lehocsky, J., Levine, R., Liu, G., Liu, G., Locke, K., Macdonald, P., Margulis, N., McCarthy, M., McEwan, P., McGurk, A., McKernan, K., McSheeters, R., Meldrum, J., Menus, L., Mihova, T., Miranda, C., Mienna, V., Morrow, J., Murphy, T., Naylor, J., Norman, C.H., O'Connor, T., O'Donnell, P., O'Neill, D., Oliver, T.M., Oliver, J., Peterson, K., Piatre, N., Pisan, C., Pollara, V., Raymond, C., Riley, R., Rogov, P., Rothman, D., Roy, A., Santos, R., Schauer, S., Severy, P., Spencer, B., Stange-Thomann, N., Stojanovic, N., Subramanian, A., Talmas, J., Tsafaye, S., Theodore, J., Tittrell, A., Travers, M., Triggiani, J., Vassiliev, H., Viel, R., Vo, A., Wilson, B., Wu, X., Wyman, D., Ye, W.J., Young, G., Zainoun, J., Zimmer, A. and Zody, M.
TITLE			Direct Submmission
JOURNAL			Submitted (12-MAY-2000) Whitehead Institute/MIT Center for Genome Research, 320 Charles Street, Cambridge, MA 02141, USA
COMMENT			On Dec 7, 2000 this sequence version replaced gi:8671969.
			All repeats were identified using RepeatMasker:
			Smt, A.F.A. & Green, P. (1996-1997)
			http://ftp.genome.washington.edu/RM/RepeatMasker.html
			Genome Center
			Center: Whitehead Institute/ MIT Center for Genome Research
			Center code: WTHR
			Web site: http://www-beg.wi.mit.edu
			Contact: sequence_submissions@genome.wi.mit.edu
			Project Information
			Center project name: L7679
			Center clone name: 206 C 14
			Summary Statistics
			Sequencing vector: M13, M7815, 42% of reads
			Sequencing vector: Plasmid, n/a, 58% of reads

FEATURES	source
misc_feature	Chemistry: Dye-terminator Big Dye; 100% of reads Assembly program: Phrap; version 0.960731 Consensus quality: 172732 bases at least Q40 Consensus quality: 175502 bases at least Q30 Consensus quality: 176567 bases at least Q20 Insert size: 170000; agarose-fp Insert size: 177079; sum-of-ctnigs Quality coverage: 6.2 in Q20 bases; agarose-fp Quality coverage: 5.9 in Q20 b. NOTE: This is a 'working draft' sequence. It currently consists of 11 contigs. The true order of the pieces is not known and their order in this sequence record is arbitrary. Gaps between the contigs are represented as runs of N, but the exact sizes of the gaps are unknown. This record will be updated with the finished sequence as soon as it is available and the accession number will be preserved.
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gap	8649 9685: contig of 1037 bp in length
misc_feature	9686 9785: gap of 100 bp
gap	9786 11207: contig of 1422 bp in length
misc_feature	11208 11307: gap of 100 bp
gap	11308 14537: contig of 3230 bp in length
misc_feature	14538 14637: gap of 100 bp
gap	14638 22686: contig of 8049 bp in length
misc_feature	22687 22786: gap of 100 bp
gap	22787 34297: contig of 1511 bp in length
misc_feature	34298 34397: gap of 100 bp
gap	34398 47850: contig of 13453 bp in length
misc_feature	47851 47950: gap of 100 bp
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gap	67381 88352: contig of 20972 bp in length
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Query Match 38.8%; Score 155.2; DB 12; Length 178079;
Best Local Similarity 73.7%; Pred. No. 6.2e-42;
Matches 196; Conservative 1; Mismatches 69; Indels 0; Gaps 0;

Qy 81 ATGCTGTAAATCCCGACGCTTCGGAGCCGAGTGGCGGATCCTTGAAGTCAAGGA 140
Db 130549 ATGCTGTAAATCCCGACGCTTCGGAGCCGAGTGGCGGATCCTTGAAGTCAAGGA 130490
Qy 141 TCGAGACCATCTGGCCCAACATGTGAACCCCGCTTAAATAAATACAAAAATAGC 200
Db 130489 TCGAGACCATCTGGCCCAACATGTGAACCCCGCTTAAATAAATACAAAAATAGC 130430
Qy 201 TGGGATGTGGGACACACCTGTAGTCCAGCTACTCAGAGCCGAGATTGACGTGAGC 260
Db 130429 CGGCTCTGTGATCTACACCTGTAGTCCAGCTACTCAGAGCCGAGATTGAGC 130370
Qy 261 TGAATTCGACAGTGAAGCCGAATTCACAGATCAGAGTGAAGCAGAGATGACACCCGCT 320
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Db 130309 CAAAAACAACAACAAAAA 130284

RESULT 49
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LOCUS AL162384
DEFINITION Human DNA sequence from clone RP11-21817 on chromosome 9 contains a novel gene similar to RAN binding protein 6 (RANBP6), the 5' end of a novel gene, a novel gene and a Cpg island, complete sequence.
ACCESSION AL162384
VERSION AL162384.14 GI:13274331
KEYWORDS HTG; Cpg island; RANBP6.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini; Homnidae; Homo.
REFERENCE 1 (bases 1 to 68465)
AUTHORS Tracey A.
TITLE Direct Submission
JOURNAL Submitted (13-MAY-2005) Wellcome Trust Sanger Institute, Hinxton, Cambridgeshire, CB10 1SA, UK. E-mail enquiries: vegas@sanger.ac.uk
Clone requests: clonerequest@sanger.ac.uk
On Mar 12, 2001 this sequence version replaced gi:12831806.
The following abbreviations are used to associate primary accession numbers given in the feature table with their source databases: Emi, ENBL; Sw, SWISSPROT; Tr, TREMBL; Wp, WORMPEP; Information on the WORMPEP database can be found at
http://www.sanger.ac.uk/Projects/C_elegans/wormpep This sequence

FEATURES

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CDS

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DEFINITION Homo sapiens 12 BAC RP11-774122 (Roswell Park Cancer Institute
ACCESSION Human BAC Library) complete sequence.
VERSION AC068889
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SOURCE HTG.
ORGANISM Homo sapiens (human)
REFERENCE
AUTHORS Murny D.M., Adams C., Adio-Oduola B., Ali-oshan F.R., Allen C.,
Albrooks S.L., Amaralunge H.C., Are J.R., Ayale M., Banks T.,
Barbata J., Benton J., Blumage K., Blankenburg K., Bonini D.,
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Weinstock G., and Gibbs R.
Direct Submission
TITLE Unpublished
AUTHORS 2 (bases 1 to 73433)
Worley K.C.
REFERENCE
AUTHORS Direct Submission
TITLE Submitted (11-MAY-2000) Human Genome Sequencing Center, Department
of Molecular and Human Genetics, Baylor College of Medicine, One
Baylor Plaza, Houston, TX 77030, USA
3 (bases 1 to 73433)
Worley K.C.
REFERENCE
AUTHORS Direct Submission
TITLE Submitted (21-SEP-2002) Human Genome Sequencing Center, Department
of Molecular and Human Genetics, Baylor College of Medicine, One
Baylor Plaza, Houston, TX 77030, USA
4 (bases 1 to 73433)
Worley K.C.
REFERENCE
AUTHORS Direct Submission
TITLE Submitted (01-JAN-2003) Human Genome Sequencing Center, Department
of Molecular and Human Genetics, Baylor College of Medicine, One
Baylor Plaza, Houston, TX 77030, USA

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RESULT 50
AC068889
LOCUS AC068889 73433 bp DNA linear PRI 01-JAN-2003

COMMENT

On Jan 1, 2003, this sequence version replaced gi:23264934.
 INFORMATION: <http://www.hgsc.bcm.tmc.edu/> or email
gc-help@bcm.tmc.edu

CLONE LENGTH: This sequence does not necessarily represent the
 entire insert of this clone. Overlapping regions of clones are only
 sequenced and submitted once, so the sequence for the remainder of
 the insert may be found in the record for the adjacent clones.
 Overlapping clones are noted at the beginning and end of the
 Features listing.

ANNOTATION OF FEATURES:

STSs are identified using ePCR (Genome Res. 7:541-550) searches
 of a local database that includes entries from dbSTS, GDB, and
 local mapping efforts.

Repeats are identified using RepeatMasker (A. Smit and P. Green,
 unpublished.) for Human and Mouse sequences.

Genes and Region of sequence similarity are identified by BLAST
 (Nuc. Acids Res. 25:3389-3402) similarity (expect < 1e-34) to the
 EST and cDNA sequences. Genes demonstrate at least two exons
 flanked by consensus splice sites that maintained sequence
 continuity across the splice junctions. Sequences that are not
 identical matches are annotated as similar.

SEQUENCING READ COVERAGE: Sequencing is completed to a minimum
 standard of double strand coverage with a minimum of 2 clones and 2
 reads with no ambiguities or 2 chemistries with a minimum of 2
 clones and 3 reads with no ambiguities. If the sequence quality for
 a region does not meet this standard, it will be indicated in the
 annotation as Low Coverage.

QUALITY OF INDIVIDUAL BASES: This sequence meets stringent quality
 standards - estimated error rate less than 1 per 10,000 bases.
 Reports of lowest quality individual bases and measures of base
 quality are listed below. Description of the metrics can be found
 at URL:
<http://www.hgsc.bcm.tmc.edu:8088/quality.info/genbank.annotation.html>.

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QY	261	TGAGATCCCAAGTAGGCGCAATCAGATCAGAGTGAAGAGAGGAGACCCGCT	320
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OY 321 CAAAACAAACAAAACAAAACCATAGA 355
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Search completed: July 17, 2006, 21:02:06
Job time : 2733 secs

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GenCore version 5.1.9
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OM nucleic - nucleic search, using sw model

Run on: July 17, 2006, 20:02:44 ; Search time 525 Seconds
(without alignments)
5312.189 Million cell updates/sec

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Perfect score: 399.6
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Scoring table: IDENTITY_NUC
Gapop 10.0 , Gapext 1.0

Searched: 5244920 seqs, 3486124231 residues
Total number of hits satisfying chosen parameters: 10489840

Minimum DB seq length: 0
Maximum DB seq length: 200000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 150 summaries

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- 3: geneeqn2000s:*
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- 10: geneeqn2003cs:*
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- 12: geneeqn2004as:*
- 13: geneeqn2004bs:*
- 14: geneeqn2005s:*
- 15: geneeqn2006s:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

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7	158.4	39.6	96593	10	ADB72488 Human TRF
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PR 14-SEP-2000; 2000US-0233065P.
PR 21-SEP-2000; 2000US-0234223P.
PR 21-SEP-2000; 2000US-0234224P.
PR 25-SEP-2000; 2000US-0234997P.
PR 25-SEP-2000; 2000US-0234998P.
PR 26-SEP-2000; 2000US-0235484P.
PR 27-SEP-2000; 2000US-0235834P.
PR 27-SEP-2000; 2000US-0235836P.
PR 29-SEP-2000; 2000US-0236327P.
PR 29-SEP-2000; 2000US-0236367P.
PR 29-SEP-2000; 2000US-0236368P.
PR 29-SEP-2000; 2000US-0236369P.
PR 29-SEP-2000; 2000US-0236370P.
PR 02-OCT-2000; 2000US-0236802P.
PR 02-OCT-2000; 2000US-0237037P.
PR 02-OCT-2000; 2000US-0237038P.
PR 02-OCT-2000; 2000US-0237039P.
PR 02-OCT-2000; 2000US-0237040P.
PR 13-OCT-2000; 2000US-0239933P.
PR 13-OCT-2000; 2000US-0239937P.
PR 20-OCT-2000; 2000US-0240960P.
PR 20-OCT-2000; 2000US-0241221P.
PR 20-OCT-2000; 2000US-0241785P.
PR 20-OCT-2000; 2000US-0241786P.
PR 20-OCT-2000; 2000US-0241787P.
PR 20-OCT-2000; 2000US-0241808P.
PR 20-OCT-2000; 2000US-0241809P.
PR 01-NOV-2000; 2000US-0241826P.
PR 01-NOV-2000; 2000US-0244617P.
PR 08-NOV-2000; 2000US-0246474P.
PR 08-NOV-2000; 2000US-0246475P.
PR 08-NOV-2000; 2000US-0246477P.
PR 08-NOV-2000; 2000US-0246478P.
PR 08-NOV-2000; 2000US-0246523P.
PR 08-NOV-2000; 2000US-0246524P.
PR 08-NOV-2000; 2000US-0246525P.
PR 08-NOV-2000; 2000US-0246526P.
PR 08-NOV-2000; 2000US-0246527P.
PR 08-NOV-2000; 2000US-0246528P.
PR 08-NOV-2000; 2000US-0246532P.
PR 08-NOV-2000; 2000US-0246609P.
PR 08-NOV-2000; 2000US-0246610P.
PR 08-NOV-2000; 2000US-0246611P.
PR 08-NOV-2000; 2000US-0246613P.
PR 17-NOV-2000; 2000US-0249207P.
PR 17-NOV-2000; 2000US-0249208P.
PR 17-NOV-2000; 2000US-0249209P.
PR 17-NOV-2000; 2000US-0249210P.
PR 17-NOV-2000; 2000US-0249211P.
PR 17-NOV-2000; 2000US-0249212P.
PR 17-NOV-2000; 2000US-0249213P.
PR 17-NOV-2000; 2000US-0249214P.
PR 17-NOV-2000; 2000US-0249215P.
PR 17-NOV-2000; 2000US-0249216P.
PR 17-NOV-2000; 2000US-0249217P.
PR 17-NOV-2000; 2000US-0249218P.
PR 17-NOV-2000; 2000US-0249244P.
PR 17-NOV-2000; 2000US-0249245P.
PR 17-NOV-2000; 2000US-0249246P.
PR 17-NOV-2000; 2000US-0249265P.
PR 17-NOV-2000; 2000US-0249297P.
PR 17-NOV-2000; 2000US-0249299P.
PR 17-NOV-2000; 2000US-0249300P.
PR 01-DEC-2000; 2000US-0250160P.
PR 01-DEC-2000; 2000US-0250391P.
PR 05-DEC-2000; 2000US-0251030P.
PR 05-DEC-2000; 2000US-0251988P.
PR 05-DEC-2000; 2000US-0256719P.
PR 06-DEC-2000; 2000US-0251479P.
PR 08-DEC-2000; 2000US-0251856P.
PR 08-DEC-2000; 2000US-0251868P.

PR 08-DEC-2000; 2000US-0251869P.
PR 08-DEC-2000; 2000US-0251989P.
PR 08-DEC-2000; 2000US-0251980P.
PR 11-DEC-2000; 2000US-0254097P.
PR 05-JAN-2001; 2001US-0259678P.
XX
XX (HUMA-) HUMAN GENOME SCI INC.
PI Rosen CA, Barash SC, Ruben SM;
XX WPI; 2001-483426/52.
XX
XX Nucleic acids encoding human immune/hematopoietic antigen polypeptides,
PT useful for preventing, diagnosing and/or treating cancers and metastasis.
XX
XX Disclosure; SEQ ID NO 38713; 3071pp + Sequence Listing; English.
XX
XX AAK54951 to AAK64702 encode the human immune/haematopoietic antigen (I)
CC amino acid sequences given in AAM82170 to AAM91921. (I) have cytosolic
CC activity, and can be used in gene therapy and vaccine production. (I)
CC proteins and polynucleotides may be used in the prevention, diagnosis and
CC treatment of diseases associated with inappropriate (I) expression. For
CC example, they may be used to treat disorders associated with decreased
CC expression by rectifying mutations or deletions in a patient's genome
CC that affect the activity of (I) by expressing inactive proteins or to
CC supplement the patient's own production of (I). Additionally, (I)
CC polynucleotides may be used to produce the secreted (I), by inserting the
CC nucleic acids into a host cell and culturing the cell to express the
CC protein. (I) proteins and polynucleotides may be used to prevent,
CC diagnose and treat immune/haematopoietic-related diseases, especially
CC cancers and cancer metastases of haematopoietic-derived cells. AAK64703
CC to AAK87694 represent human immune/haematopoietic antigen genomic
CC sequences from the present invention. AAK54942 to AAK54950 and AAM82169
CC represent sequences used in the exemplification of the present invention
XX
XX SQ Sequence 16163 BP; 4414 A; 3239 C; 3448 G; 5062 T; 0 U; 0 Other;
XX
XX Query Match 39.6%; Score 158.4; DB 4; Length 16163;
XX Best Local Similarity 73.9%; Pred. No. 1e-33;
XX Matches 201; Conservative 0; Mismatches 71; Indels 0; Gaps 0;
QY 76 GCTGAATGCTCTTAATCCGACCTTCGGAGGCGCAAGTGGCGGATCACTGAGTCA 135
DB 7992 GATTACGCGCTGTATCCGACACTCTGGAGGCGCGGTGTATCACTATATCA 7933
QY 136 AGAATGAGACCATCTCTGGCCATGCTGTAACCCCGTCTTACTAAATCAAAAA 195
DB 7932 GAGATTGAGATCAGTCTGGCCACATGTGTAACCCCGTCTTACTAAATCAAAAA 7873
QY 196 ATAGCTGAGCATGTGTCACACACTGTAGTCCAGCTACTCAGAGCCGGAGATTGCAG 255
DB 7872 CCACTGGGTGTGTGTGTAACCGTTGTATCCAGTACTCTGAGGCGGAGTTGCAG 7813
QY 256 TGAGCTGAGATCGCAGAGTGGCCGAATCATCAGATCAAGAGTGCAGCATGACAC 315
DB 7812 TGAGCCGAAACTGACACATCACTCCAGCTGGGTGACAGAGGAGACTCTGTCCCA 7753
QY 316 CGTCTCAAAAACACACACAAAAACAAAAA 347
DB 7752 GGAATAAAAAAAAAAAAAAAAAAAAAA 7721

RESULT 4
ID AAK87161/C
ID AAK87161 standard; DNA; 16163 BP.
XX
XX AAK87161;
AC
AC 07-NOV-2001 (first entry)
XX
XX Human immune/haematopoietic antigen genomic sequence SEQ ID NO:41973.
DE Human immune/haematopoietic; immune/haematopoietic antigen; cancer;
XX
XX Human; immune; haematopoietic; immune/haematopoietic antigen; cancer;
```

KY cytostatic; gene therapy; vaccine; metastasis; ds.
XX Homo sapiens.
XX WO200157182-A2.
XX 09-AUG-2001.
XX 17-JAN-2001; 2001WO-US001354.
XX 31-JAN-2000; 2000US-0179065P.
XX 04-FEB-2000; 2000US-0180628P.
XX 24-FEB-2000; 2000US-0184664P.
XX 02-MAR-2000; 2000US-0186350P.
XX 16-MAR-2000; 2000US-0188874P.
XX 17-MAR-2000; 2000US-0190076P.
XX 18-APR-2000; 2000US-0198123P.
XX 19-MAY-2000; 2000US-0205151P.
XX 07-JUN-2000; 2000US-0209467P.
XX 28-JUN-2000; 2000US-0214886P.
XX 30-JUN-2000; 2000US-0215135P.
XX 07-JUL-2000; 2000US-0216647P.
XX 07-JUL-2000; 2000US-0216880P.
XX 11-JUL-2000; 2000US-0217487P.
XX 11-JUL-2000; 2000US-0217496P.
XX 14-JUL-2000; 2000US-0218290P.
XX 26-JUL-2000; 2000US-0220963P.
XX 26-JUL-2000; 2000US-0220966P.
XX 14-AUG-2000; 2000US-0224518P.
XX 14-AUG-2000; 2000US-0224519P.
XX 14-AUG-2000; 2000US-0225213P.
XX 14-AUG-2000; 2000US-0225214P.
XX 14-AUG-2000; 2000US-0225266P.
XX 14-AUG-2000; 2000US-0225267P.
XX 14-AUG-2000; 2000US-0225268P.
XX 14-AUG-2000; 2000US-0225270P.
XX 14-AUG-2000; 2000US-0225447P.
XX 14-AUG-2000; 2000US-0225757P.
XX 14-AUG-2000; 2000US-0225758P.
XX 14-AUG-2000; 2000US-0225759P.
XX 18-AUG-2000; 2000US-0226279P.
XX 22-AUG-2000; 2000US-0226681P.
XX 22-AUG-2000; 2000US-0226686P.
XX 22-AUG-2000; 2000US-0227182P.
XX 23-AUG-2000; 2000US-0227109P.
XX 30-AUG-2000; 2000US-0228924P.
XX 01-SEP-2000; 2000US-0229287P.
XX 01-SEP-2000; 2000US-0229343P.
XX 01-SEP-2000; 2000US-0229344P.
XX 01-SEP-2000; 2000US-0229345P.
XX 05-SEP-2000; 2000US-0229509P.
XX 05-SEP-2000; 2000US-0229513P.
XX 06-SEP-2000; 2000US-0230437P.
XX 06-SEP-2000; 2000US-0230438P.
XX 08-SEP-2000; 2000US-0231242P.
XX 08-SEP-2000; 2000US-0231243P.
XX 08-SEP-2000; 2000US-0231244P.
XX 08-SEP-2000; 2000US-0231244P.
XX 08-SEP-2000; 2000US-0231411P.
XX 08-SEP-2000; 2000US-0231414P.
XX 08-SEP-2000; 2000US-0232080P.
XX 08-SEP-2000; 2000US-0232081P.
XX 12-SEP-2000; 2000US-0231968P.
XX 14-SEP-2000; 2000US-0232397P.
XX 14-SEP-2000; 2000US-0233398P.
XX 14-SEP-2000; 2000US-0233399P.
XX 14-SEP-2000; 2000US-0233400P.
XX 14-SEP-2000; 2000US-0233401P.
XX 14-SEP-2000; 2000US-0233063P.
XX 14-SEP-2000; 2000US-0233064P.
XX 14-SEP-2000; 2000US-0233065P.
XX 21-SEP-2000; 2000US-0234223P.
XX 21-SEP-2000; 2000US-0234274P.
XX 25-SEP-2000; 2000US-0234997P.

PR 25-SEP-2000; 2000US-0234998P.
PR 26-SEP-2000; 2000US-0235484P.
PR 27-SEP-2000; 2000US-0235834P.
PR 27-SEP-2000; 2000US-0235836P.
PR 29-SEP-2000; 2000US-0236327P.
PR 29-SEP-2000; 2000US-0236367P.
PR 29-SEP-2000; 2000US-0236368P.
PR 29-SEP-2000; 2000US-0236369P.
PR 29-SEP-2000; 2000US-0236370P.
PR 02-OCT-2000; 2000US-0236802P.
PR 02-OCT-2000; 2000US-0237037P.
PR 02-OCT-2000; 2000US-0237038P.
PR 02-OCT-2000; 2000US-0237039P.
PR 02-OCT-2000; 2000US-0237040P.
PR 13-OCT-2000; 2000US-0239935P.
PR 13-OCT-2000; 2000US-0239937P.
PR 20-OCT-2000; 2000US-0240960P.
PR 20-OCT-2000; 2000US-0241221P.
PR 20-OCT-2000; 2000US-0241785P.
PR 20-OCT-2000; 2000US-0241786P.
PR 20-OCT-2000; 2000US-0241787P.
PR 20-OCT-2000; 2000US-0241808P.
PR 20-OCT-2000; 2000US-0241809P.
PR 01-NOV-2000; 2000US-0244617P.
PR 08-NOV-2000; 2000US-0246474P.
PR 08-NOV-2000; 2000US-0246475P.
PR 08-NOV-2000; 2000US-0246476P.
PR 08-NOV-2000; 2000US-0246477P.
PR 08-NOV-2000; 2000US-0246478P.
PR 08-NOV-2000; 2000US-0246523P.
PR 08-NOV-2000; 2000US-0246524P.
PR 08-NOV-2000; 2000US-0246525P.
PR 08-NOV-2000; 2000US-0246526P.
PR 08-NOV-2000; 2000US-0246527P.
PR 08-NOV-2000; 2000US-0246528P.
PR 08-NOV-2000; 2000US-0246529P.
PR 08-NOV-2000; 2000US-0246609P.
PR 08-NOV-2000; 2000US-0246610P.
PR 08-NOV-2000; 2000US-0246611P.
PR 08-NOV-2000; 2000US-0246613P.
PR 17-NOV-2000; 2000US-0249207P.
PR 17-NOV-2000; 2000US-0249208P.
PR 17-NOV-2000; 2000US-0249209P.
PR 17-NOV-2000; 2000US-0249210P.
PR 17-NOV-2000; 2000US-0249211P.
PR 17-NOV-2000; 2000US-0249212P.
PR 17-NOV-2000; 2000US-0249213P.
PR 17-NOV-2000; 2000US-0249214P.
PR 17-NOV-2000; 2000US-0249215P.
PR 17-NOV-2000; 2000US-0249216P.
PR 17-NOV-2000; 2000US-0249217P.
PR 17-NOV-2000; 2000US-0249218P.
PR 17-NOV-2000; 2000US-0249244P.
PR 17-NOV-2000; 2000US-0249245P.
PR 17-NOV-2000; 2000US-0249246P.
PR 17-NOV-2000; 2000US-0249265P.
PR 17-NOV-2000; 2000US-0249297P.
PR 17-NOV-2000; 2000US-0249299P.
PR 17-NOV-2000; 2000US-0249300P.
PR 01-DEC-2000; 2000US-0250160P.
PR 01-DEC-2000; 2000US-0250311P.
PR 05-DEC-2000; 2000US-0251030P.
PR 05-DEC-2000; 2000US-0251988P.
PR 05-DEC-2000; 2000US-0256719P.
PR 06-DEC-2000; 2000US-0251479P.
PR 08-DEC-2000; 2000US-0251856P.
PR 08-DEC-2000; 2000US-0251856P.
PR 08-DEC-2000; 2000US-0251869P.
PR 08-DEC-2000; 2000US-0251989P.
PR 11-DEC-2000; 2000US-0254097P.
PR 05-JAN-2001; 2001US-0259678P.

(HUMA-) HUMAN GENOME SCI INC.
Rosen CA, Barash SC, Ruben SM;

PT New recombinant nucleic acid encoding carcinoma associated protein, useful for immunizing animals to prevent cancer

XX

PS Claim 1; SEQ ID NO 1268; 245bp; English.
XX
CC The invention relates to recombinant carcinoma associated (CA) nucleic
CC acid sequences from mouse and human (AD801482-ADA03094), and to
CC recombinant carcinoma associated proteins (CAP) encoded by them. The
CC invention also encompasses expression vectors and host cells comprising a
CC CA nucleic acid, a polypeptide (especially an antibody) that specifically
CC binds to the protein, and a bioclip comprising CA nucleic acid or
CC fragments thereof. The sequences of the invention were identified using
CC oncogenic retroviruses, which insert into the genome of the host organism
CC at random. Many of these do not carry transduced host oncogenes or
CC pathogenic trans-acting viral genes, meaning that cancer incidence is a
CC direct consequence of the effects of proviral integration into host
CC protooncogenes. The CA nucleic acid sequences can be used to diagnose
CC carcinoma (especially breast cancer, prostate cancer, lymphoma or
CC leukemia) or a propensity to carcinoma by determination of the sequence
CC of a CA gene, or by determination of CA gene expression in particular
CC tissues. CA nucleic acids, proteins and antibodies are also useful as
CC therapeutic agents and in screening and evaluating drug candidates. The
CC present sequence represents a specifically claimed human CA nucleic acid
CC sequence of the invention. Note: The complete sequence data for this
CC patent did not form part of the printed specification, but was obtained
CC in electronic format directly from WIPO at
CC ftp.wipo.int/pub/published_pct_sequences.
XX
SQ Sequence 96593 BP; 25039 A; 20903 C; 22979 G; 27672 T; 0 U; 0 Other;

Query Match 39.6%; Score 158.4; DB 9; Length 96593;
Best Local Similarity 75.0%; Pred. No. 1.8e-33;
Matches 216; Conservative 1; Mismatches 57; Indels 14; Gaps 1;

QY 81 ATGCGCTTAATCCAGACCTTGGGAGGCCAAGGTGGCGGATCACCTGAGTCAAGAGA 140
DB 18217 ACGGCTGTAATCCAGACACTTTGGAGGCCGAGCGCGGATTTCTTGAGGTCAAGAGT 18276
QY 141 TCGAGACCATCTTGCGCAACATGTGTAACCCCGTCTTTACTATAAAATACAAAAATAGC 200
DB 18277 TCAAGATCAGCTTGCGCAACATGTGTAACCCCGTCTTTACTATAAAATACAAAAATAGC 18336
QY 201 TGGGCATGTGTGGCAACACCTGTAGTCCAGCTACTCAGAGCGCGAGATTGCAGTAGC 260
DB 18337 TGGGCATGTGTGGCGGACCTGTAGTCCAGCTACTCAGAGCGCGAGAAATCG 18396
QY 261 TGAATGCGCAG-----AGTGAGCCGGAATTCACAGATCACAGAGTAGAGCAGA 306
DB 18397 CTTGAATCCAGGCGCGGAGGTTGCAGTGAAGCCGAGATCGACAGCTTGCGTGAAGC 18456
QY 307 GTGAGACKCCGCTCTCAAAAACACAAACAAAAAACAATTAAG 354
DB 18457 GTGAGACTTCATCTCAAAAAAACAACAAAAACAACCAACGATG 18504

RESULT 7
ADB72488
ID ADB72488 standard; DNA; 96593 BP.
AC ADB72488;
XX
XX 04-DEC-2003 (first entry)
XX
XX Human IRF2 gene.
XX
XX human; ds; cytosratic; gene therapy; vaccine; carcinoma; lymphomas;
XX cancer; neoplasm; adenocarcinoma; sarcoma; gene.
XX
XX Homo sapiens.
XX
XX MO2003008583-A2.
XX
XX 30-JAN-2003.
XX
XX 26-DEC-2001; 2001WO-US051291.
XX

PR 02-MAR-2001; 2001US-00798586.
PR 23-OCT-2001; 2001US-00004113.
PR 08-NOV-2001; 2001US-00002482.
PR 30-NOV-2001; 2001US-00997722.
PR 20-DEC-2001; 2001US-00034650.
XX
XX (SAGR-) SAGRES DISCOVERY.
XX
XX Morris DW, Engelhard EK;
XX WPI; 2003-229337/23.
XX
XX New recombinant nucleic acid, useful for treating carcinomas, lymphomas,
XX cancers, neoplasm, adenocarcinoma, or sarcomas.
XX
XX
XX Claim 1; SEQ ID NO 316; 2304pp; English.
XX
XX The invention relates to a novel recombinant nucleic acid comprising a
XX nucleotide sequence selected from any of the 660 sequences fully defined
XX in the specification. A polynucleotide of the invention has cytosratic
XX activity and may have a use in gene therapy, or in a vaccine. The
XX recombinant nucleic acids and polypeptides are useful for treating
XX carcinomas, e.g. lymphomas, cancers, neoplasm, adenocarcinoma, and
XX sarcomas. The present sequence represents a human gene of the invention.
XX
SQ Sequence 96593 BP; 25039 A; 20903 C; 22979 G; 27672 T; 0 U; 0 Other;

Query Match 39.6%; Score 158.4; DB 10; Length 96593;
Best Local Similarity 75.0%; Pred. No. 1.8e-33;
Matches 216; Conservative 1; Mismatches 57; Indels 14; Gaps 1;

QY 81 ATGCGCTTAATCCAGACCTTGGGAGGCCAAGGTGGCGGATCACCTGAGTCAAGAGA 140
DB 18217 ACGGCTGTAATCCAGACACTTTGGAGGCCGAGCGCGGATTTCTTGAGGTCAAGAGT 18276
QY 141 TCGAGACCATCTTGCGCAACATGTGTAACCCCGTCTTTACTATAAAATACAAAAATAGC 200
DB 18277 TCAAGATCAGCTTGCGCAACATGTGTAACCCCGTCTTTACTATAAAATACAAAAATAGC 18336
QY 201 TGGGCATGTGTGGCAACACCTGTAGTCCAGCTACTCAGAGCGCGGAGATTGCAGTAGC 260
DB 18337 TGGGCATGTGTGGCGGACCTGTAGTCCAGCTACTCAGAGCGCGAGAAATCG 18396
QY 261 TGAATGCGCAG-----AGTGAGCCGGAATTCACAGATCACAGAGTAGAGCAGA 306
DB 18397 CTTGAATCCAGGCGCGGAGGTTGCAGTGAAGCCGAGATCGACAGCTTGCGTGAAGC 18456
QY 307 GTGAGACKCCGCTCTCAAAAACACAAACAAAAAACAATTAAG 354
DB 18457 GTGAGACTTCATCTCAAAAAAACAACAAAAACAACCAACGATG 18504

RESULT 8
ADC85230
ID ADC85230 standard; DNA; 96593 BP.
XX
XX ADC85230;
XX
XX 01-JAN-2004 (first entry)
XX
XX Human Irf2 genomic sequence.
XX
XX
XX Cytosratic; gene therapy; vaccine; cancer; carcinoma-associated gene; CA;
XX secreted; transmembrane; intracellular; ds.
XX
XX Homo sapiens.
XX
XX MO2003045230-A2.
XX
XX 05-JUN-2003.
XX
XX 02-DEC-2002; 2002WO-US038582.
XX

PR 30-NOV-2001; 2001US-00997722.
 XX (SAGR-) SAGRES DISCOVERY.
 XX
 XX Morris DW, Engelhard EK;
 XX WPI; 2003-513603/48.
 XX
 PT New recombinant nucleic acid comprising a nucleotide sequence of any of
 PT the carcinoma-associated (CA) genes, useful for screening for drug
 PT candidates for diagnosing or treating carcinomas.
 XX
 PS Claim 1; SEQ ID NO 16; 983pp; English.
 XX
 CC The invention relates to a recombinant nucleic acid comprising a
 CC nucleotide sequence selected from any of the fully defined carcinoma-
 CC associated (CA) genes from the 50 tables given in the specification. The
 CC CA proteins are secreted, transmembrane or intracellular proteins. The
 CC recombinant nucleic acids are useful for screening for drug candidates
 CC for diagnosing or treating carcinomas. Sequences given in A0685215-
 CC A0685514 represent CA genes of the invention.
 XX
 SQ Sequence 96593 BP; 25039 A; 20903 C; 22979 G; 27672 T; 0 U; 0 Other;
 Query Match 39.6%; Score 158.4; DB 10; Length 96593;
 Best Local Similarity 75.0%; Pred. No. 1.8e-33;
 Matches 216; Conservative 1; Mismatches 57; Indels 14; Gaps 1;
 QY 81 ATGCTGTATTCAGCACTTCGGAGGCGCAAGTGGGCGGATCACTGAGGTCAAGAGA 140
 DB 18217 ACGCTGTATTCAGCACTTCGGAGGCGGAGGCGGAGATTCTGAGGTCAAGAGT 18276
 QY 141 TCGAGACCATCTGGCCCAACATGTGAAACCCCGCTTTTACTTAAATAACAAAAATAGC 200
 DB 18277 TCAAGATAGCTTGGCCCAACATGTGAAACCCCGCTTTTACTTAAATAACAAAAATAGC 18336
 QY 201 TGGGATGTGGACACACCTGTATGTCAGCTACTCGAGAGCGGAGATTCAGATGAGC 260
 DB 18337 TGGGATGTGGGCGGACCTGTATGTCAGCTACTCGAGAGCGGAGATTCAGATGAGC 18396
 QY 261 TGAATGCGAG-----AGTAGCCGAATTCACAGATCAGAGTGAAGCAGA 306
 DB 18397 CTTGAATGCGAGCGGCGGAGGTTGCAAGAGCGGAGATTCAGATGAGTGAAGCAGA 18456
 QY 307 GTGAGACCCGCTCTCAAAAACAACAACAACAAAAAACCAATAG 354
 DB 18457 GTGAGACTTCATCTCAAAAACAACAACAACAAAAACAACCAAGATG 18504
 RESULT 9
 ADM74345
 ID ADM74345 standard; DNA; 96593 BP.
 XX
 AG ADM74345;
 XX
 DT 01-JUL-2004 (first entry)
 XX
 DB Human carcinoma associated (CA) nucleic acid #7.
 XX
 XX Human; carcinoma associated nucleic acid; CA nucleic acid; gene; de;
 XX carcinoma associated protein; CAP; carcinoma; leukemia; lymphoma;
 XX cytosolic.
 XX
 OS Homo sapiens.
 XX
 XX US2004072154-A1.
 XX
 XX 15-APR-2004.
 XX
 XX 30-NOV-2001; 2001US-00997722.
 XX
 XX 22-DEC-2000; 2000US-00747377.
 XX
 XX 02-MAR-2001; 2001US-00798586.
 PR

XX
 PA (MORRIS) MORRIS D W.
 PA (ENGELHARD) ENGELHARD E K.
 XX
 XX Morris DW, Engelhard EK;
 XX WPI; 2004-328562/30.
 XX
 PT New carcinoma associated gene or protein, useful for preparing a
 PT composition for diagnosing or treating carcinoma e.g., leukemia or
 PT lymphoma.
 XX
 PS Claim 1; SEQ ID NO 16; 29pp; English.
 XX
 CC The invention relates to new recombinant nucleic acid. The invention
 CC also relates to a host cell comprising a recombinant nucleic acid or
 CC expression vector, an expression vector comprising a recombinant nucleic
 CC acid, a recombinant protein, a method of screening for drug candidates, a
 CC method of screening for a bioactive agent capable of binding to a
 CC carcinoma associated protein (CAP) encoded by a nucleotide sequence, a
 CC method of screening for a bioactive agent capable of modulating the
 CC activity of a CAP, a method of evaluating the effect of a candidate
 CC carcinoma drug, a method of diagnosing carcinoma, a method for inhibiting
 CC the activity of a CAP, a method of treating carcinomas, a method of
 CC neutralising the effect of a CAP and a method of diagnosing carcinoma or
 CC propensity to carcinoma. A method of evaluating the effect of a candidate
 CC carcinoma drug comprises administering the drug to a patient, removing a
 CC cell sample from the patient and determining alterations in the
 CC expression or activation of a gene comprising the nucleotide sequence. A
 CC method of diagnosing carcinoma comprises determining the expression of
 CC one or more genes comprising the nucleic acid sequence in a first tissue
 CC type of a first individual and comparing the expression of the gene from
 CC a second normal tissue type from the first individual or a second
 CC unaffected individual, where a difference in the expression indicates
 CC that the first individual has carcinoma. A method of inhibiting the
 CC activity of a CAP comprises binding an inhibitor to the CAP. Treating
 CC carcinomas comprises administering to a patient an inhibitor of CAP.
 CC Neutralising the effect of a CAP comprises contacting an agent specific
 CC for the CAP. The polypeptide specifically binds to the protein encoded by
 CC the nucleic acid. It comprises an antibody that specifically binds to the
 CC protein encoded by the nucleic acid. The nucleic acids are useful for
 CC preparing a composition for diagnosing or treating carcinoma e.g.,
 CC leukemia or lymphoma. This sequence represents a human carcinoma
 CC associated (CA) nucleic acid of the invention. Note: The sequence data
 CC for this patent did not form part of the printed specification but was
 CC obtained in electronic format directly from USPTO at
 CC seqdata.uspto.gov/sequence.html.
 XX
 SQ Sequence 96593 BP; 25039 A; 20903 C; 22979 G; 27672 T; 0 U; 0 Other;
 Query Match 39.6%; Score 158.4; DB 12; Length 96593;
 Best Local Similarity 75.0%; Pred. No. 1.8e-33;
 Matches 216; Conservative 1; Mismatches 57; Indels 14; Gaps 1;
 QY 81 ATGCTGTATTCAGCACTTCGGAGGCGCAAGTGGGCGGATCACTGAGGTCAAGAGA 140
 DB 18217 ACGCTGTATTCAGCACTTCGGAGGCGGAGGCGGAGATTCTGAGGTCAAGAGT 18276
 QY 141 TCGAGACCATCTGGCCCAACATGTGAAACCCCGCTTTTACTTAAATAACAAAAATAGC 200
 DB 18277 TCAAGATAGCTTGGCCCAACATGTGAAACCCCGCTTTTACTTAAATAACAAAAATAGC 18336
 QY 201 TGGGATGTGGACACACCTGTATGTCAGCTACTCGAGAGCGGAGATTCAGATGAGC 260
 DB 18337 TGGGATGTGGGCGGACCTGTATGTCAGCTACTCGAGAGCGGAGATTCAGATGAGC 18396
 QY 261 TGAATGCGAG-----AGTAGCCGAATTCACAGATCAGAGTGAAGCAGA 306
 DB 18397 CTTGAATGCGAGCGGCGGAGGTTGCAAGAGCGGAGATTCAGATGAGTGAAGCAGA 18456
 QY 307 GTGAGACCCGCTCTCAAAAACAACAACAACAAAAAACCAATAG 354
 DB 18457 GTGAGACTTCATCTCAAAAACAACAACAACAAAAACAACCAAGATG 18504

RESULT 10
ACN4354/c
ID ACN44354 standard; DNA; 31898 BP.
XX
AC ACN44354;
XX
DT 18-NOV-2004 (first entry)
XX
DE Human genomic sequence hCG17346.
XX
KM Cytostatic; carcinoma; lymphoma; cancer; human; gene; ss.
XX
OS Homo sapiens.
XX
PN W02003073826-A2.
XX
PD 12-SEP-2003.
XX
PF 28-FEB-2003; 2003WO-US006235.
XX
PR 01-MAR-2002; 2002US-00087192.
XX
PA (SAGR-) SAGRES DISCOVERY.
XX
PI Morris DW;
XX
DR WPI; 2003-328604/31.
XX
PT Recombinant nucleic acid useful for diagnosis and treatment of carcinoma
XX comprises a nucleotide sequence.
XX
PS Claim 1; SEQ ID NO 760; OPB; English.
XX
XX The present invention relates to novel DNA and protein sequences which
XX are associated with carcinomas. The sequences are useful for: (i) for
XX screening drug candidates; (ii) for screening of bioactive agent capable
XX of binding to Carcinoma Associated Protein (CAP); (iii) for screening of
XX a bioactive agent capable of modulating the activity of CAP; (iv) for
XX evaluating the effect of a candidate carcinoma drug; (v) for treating
XX carcinoma; (vi) for inhibiting the activity of CAP; (ix) as a biochip;
XX (x) for diagnosing carcinoma or a propensity to carcinoma; and (xi) for
XX determining carcinoma associated (CA) gene copy number. In addition, the
XX CA genes are useful as DNA vaccines and the CAP are useful as markers of
XX carcinoma including lymphoma. The present sequence is one such CA coding
XX sequence. Note: This patent is an equivalent to basic patent
XX US2002182586A1, for which no sequence data was published
XX
SQ Sequence 31898 BP; 7315 A; 9118 C; 8770 G; 6695 T; 0 U; 0 Other;
Query Match 38.9%; Score 155.4; DB 11; Length 31898;
Best Local Similarity 72.6%; Pred. No. 8.8e-33;
Matches 217; Conservative 1; Mismatches 72; Indels 9; Gaps 1;
QY 81 ATGCTGTAAATCCCAAGCACTTCGGAGGCCCAAGTGGGCGGATCCCTGAGTCAAGGA 140
DB 30243 ACGCTTAATATCCCAAGCACTTGGAAAGGCCAAGGTGGGATCACTTGAAGCCAGGAAT 30184
QY 141 TCGAGACCATCTGGGCCAATGTGTAACCCCGCTTACTTAATAAATACAAAAATAGC 200
DB 30183 TCGAGACCATCTGGGCCAATGTGTAACCCCGCTTACTTAATAAATACAAAAATAGC 30124
QY 201 TGGGCGATGGTGGACACACACTGTAGTCCAGTACTCAGAGCCGAGAGATTGCAATGAGC 260
DB 30123 CCGGAGTGTGTGGGCGCTGTAGTCCCACTACTCGGAGGCGAGAGTTGCAATGAGC 30064
QY 261 TGAATATCCGACAGTAGGCGGAAATACAGATACAGAGTAGGACAGAGAGACACGCCGCT 320
DB 30063 CGAGACCAAGCGCTTGAAGCTCCAGCTGGGCGA-----CAGAGCGAGAGCTTGTCT 30013
QY 321 CAAAAACAAACAAAAAACAACATTAAGATTTGCATCTCGGTTCCAGA 379

Db 30012 CAACAACATATACAAAAAACAAGTCAAGTCTTCTGCTTGTGAGTTGGGCTCAGA 29954
|||||
RESULT 11
ACN37240
ID ACN37240 standard; DNA; 34796 BP.
XX
AC ACN37240;
XX
DT 18-NOV-2004 (first entry)
XX
DE Human periodontal disease related gene P10D SEQ ID NO:150.
XX
KM periodontal disease; polymorphism; ds; human; gene; SNP;
XX single nucleotide polymorphism.
XX
OS Homo sapiens.
XX
FH Key
FT misc_feature
Location/Qualifiers
869
/*tag= a
/standard_name= "Single nucleotide polymorphism"
/note= "Variable nucleotide T,C"
1048
/*tag= b
/standard_name= "Single nucleotide polymorphism"
/note= "Variable nucleotide T,C"
1103
/*tag= c
/standard_name= "Single nucleotide polymorphism"
/note= "Variable nucleotide G,A"
5092
/*tag= d
/standard_name= "Single nucleotide polymorphism"
/note= "Variable nucleotide A,G"
5333
/*tag= e
/standard_name= "Single nucleotide polymorphism"
/note= "Variable nucleotide C,G"
5795
/*tag= f
/standard_name= "Single nucleotide polymorphism"
/note= "Variable nucleotide T,C"
6328
/*tag= g
/standard_name= "Single nucleotide polymorphism"
/note= "Variable nucleotide A,G"
6455
/*tag= h
/standard_name= "Single nucleotide polymorphism"
/note= "Variable nucleotide A,G"
6489
/*tag= i
/standard_name= "Single nucleotide polymorphism"
/note= "Variable nucleotide C,G"
6616
/*tag= j
/standard_name= "Single nucleotide polymorphism"
/note= "Variable nucleotide A,G"
8061
/*tag= k
/standard_name= "Single nucleotide polymorphism"
/note= "Variable nucleotide G,T"
10001
/*tag= l
/standard_name= "Single nucleotide polymorphism"
/note= "Variable nucleotide A,G"
12618
/*tag= m
/standard_name= "Single nucleotide polymorphism"
/note= "Variable nucleotide A,G"
12746
misc_feature

F1	/+tag= n	/standard_name= "Single nucleotide polymorphism"
F1	/note= "Variable nucleotide A,G"	
F1	15982	
F1	/+tag= o	/standard_name= "Single nucleotide polymorphism"
F1	/note= "Variable nucleotide C,T"	
F1	15970	
F1	/+tag= p	/standard_name= "Single nucleotide polymorphism"
F1	/note= "Variable nucleotide C,G"	
F1	15961	
F1	/+tag= q	/standard_name= "Single nucleotide polymorphism"
F1	/note= "Variable nucleotide C,T"	
F1	17746	
F1	/+tag= r	/standard_name= "Single nucleotide polymorphism"
F1	/note= "Variable nucleotide A,G"	
F1	17751	
F1	/+tag= s	/standard_name= "Single nucleotide polymorphism"
F1	/note= "Variable nucleotide C,G"	
F1	18004	
F1	/+tag= t	/standard_name= "Single nucleotide polymorphism"
F1	/note= "Variable nucleotide A,G"	
F1	22945	
F1	/+tag= u	/standard_name= "Single nucleotide polymorphism"
F1	/note= "Variable nucleotide C,T"	
F1	24796	
F1	/+tag= v	/standard_name= "Single nucleotide polymorphism"
F1	/note= "Variable nucleotide C,T"	
F1	29622	
F1	/+tag= w	/standard_name= "Single nucleotide polymorphism"
F1	/note= "Variable nucleotide G,A"	
F1	29877	
F1	/+tag= x	/standard_name= "Single nucleotide polymorphism"
F1	/note= "Variable nucleotide deletion, CT"	
F1	30580	
F1	/+tag= y	/standard_name= "Single nucleotide polymorphism"
F1	/note= "Variable nucleotide G,T"	
F1	31500	
F1	/+tag= z	/standard_name= "Single nucleotide polymorphism"
F1	/note= "Variable nucleotide G,A"	
F1	32397	
F1	/+tag= aa	/standard_name= "Single nucleotide polymorphism"
F1	/note= "Variable nucleotide C,T"	
F1	32406	
F1	/+tag= ab	/standard_name= "Single nucleotide polymorphism"
F1	/note= "Variable nucleotide C,G"	
F1	32528	
F1	/+tag= ac	/standard_name= "Single nucleotide polymorphism"
F1	/note= "Variable nucleotide C,G"	
F1	33004	
F1	/+tag= ad	/standard_name= "Single nucleotide polymorphism"
F1	/note= "Variable nucleotide C,G"	
F1	33056	
F1	/+tag= ae	/standard_name= "Single nucleotide polymorphism"
F1	/note= "Variable nucleotide C,T"	
F1	33071	
F1	/+tag= af	

FT	misc_feature	/standard_name= "Single nucleotide polymorphism"
FT	misc_feature	/note= "Variable nucleotide G,T"
FT	misc_feature	/tag= ag
FT	misc_feature	/standard_name= "Single nucleotide polymorphism"
FT	misc_feature	/note= "Variable nucleotide A,C"
PN	WO2004042054-A1.	
XX		
PD	21-MAY-2004.	
XX		
PF	22-OCT-2003; 2003WO-IB004669.	
XX		
PR	23-OCT-2002; 2002JP-00308634.	
XX		
PA	(HUBI-) HUBIT GENOMIX INC.	
XX	(KAMO/) KAMO I K.	
PI	Kamoi K, Suzuki A, Numabe Y, Ji G, Muramatsu M, Baba M,	
XX		
DR	WPI, 2004-400678/37.	
XX		
PT	Single nucleotide polymorphisms associated with periodontal disease for	
XX	examination and assessment of susceptibility to periodontal disease.	
XX		
PS	Claim 9; SEQ ID NO 150; opp; Japanese.	
XX		
CC	The invention relates to a novel method for examination of periodontal	
XX	disease in which genetic polymorphisms are detected in one or more of 51	
CC	genes. The method is useful for examination, diagnosis and assessment of	
XX	periodontal disease or risk of periodontal disease and the risk of its	
CC	progression to severe, aggressive and chronic periodontal disease. The	
XX	present sequence represents a polymorphic gene of the invention	
CC		
XX		
SQ	Sequence 34796 BP; 7631 A; 9152 C; 9368 G; 8645 T; 0 U; 0 Other;	
XX		
Query Match	38.6%; Score 154.4; DB 13; Length 34796;	
XX	Best Local Similarity 72.5%; Pred. No. 1.7e-32;	
Matches 200; Conservative 0; Mismatches 76; Indels 0; Gaps 0;		
QY	81 ATGCTCTGTAATCCCAAGCACTTCGGGAGGCGCAAGGTGGCGGATCACTGAGTCAAGGA 140	
DB	4654 ATGCTCTGTAATCCCGGCACTTGGGAGGCGCAAGCGTGTGATCATTTGGCGTCAGGAGT 4713	
QY	141 TCGAGACCATCTGGCCAAACATGGTGAACCCCGCTTACTTAAATAATACAAAATAATAGC 200	
DB	4714 TTGGAGCAGCGCTGGCCAAATGATGAAGAACCCCGCTTATTAAATAATACAAAATTAGC 4773	
QY	201 TGGGCACTGTGGGACACACCTGTAGTCCCACTACTCAGAGCCGAGATTGCAAGTGC 260	
DB	4774 TGGGCACTGTGGGAGTGGCTGTAGTCCCACTACTCAGAGGCGGAGGTTGCAAGTGC 4833	
QY	261 TGAAGTCGCAAGTGAAGCCGAATTCACAGATCACAGAGTGAAGAGTGAAGACCCGCTCT 320	
DB	4834 TGAATATTGTGCGCACTTCAGCTTCAGCTTCGGTGAAGAGAGTGAAGTGTCTCAAAAAA 4893	
QY	321 CAAAAACAACAACAAAAACAAAAACCAATTAAC 356	
DB	4894 AAAAAACAAAAACAACAACAAAAACAAAAACAAAC 4929	
RESULT 12		
ID	ABT10719/c	
XX	ABT10719 standard; cDNA; 122748 BP.	
XX	ABT10719;	
AC		
XX	04-DEC-2002 (first entry)	
XX		
DE	Human breast cancer associated coding sequence SEQ ID NO: 853.	
XX		
XX	Human, breast specific gene; breast cancer; differential expression;	
XX	cytostatic; gene therapy; gene; ss	
XX		

XX OS Homo sapiens.
XX PN WO200259271-A2.
XX PD 01-AUG-2002.
XX PF 25-JAN-2002; 2002WO-US0002176.
XX PR 25-JAN-2001; 2001US-0263757P.
XX PR 25-APR-2001; 2001US-0286090P.
XX PR 23-MAY-2001; 2001US-0292517P.
XX PA (GENE-) GENE LOGIC INC.
XX PI Orr MS, Nation M, Digians JC, Zeng W;
XX DR WPI; 2002-674803/72.
XX PT Diagnosing breast cancer in a patient comprises detecting the level of
XX PT gene expression in cell or tissue samples, where a differential gene
XX PT expression is indicative of breast cancer.
XX PS Claim 1; SEQ ID NO 853; 260bp + Sequence listing; English.
XX CC The present invention relates to methods of diagnosing breast cancer in a
XX CC patient, which comprise detecting the level of expression in a tissue
XX CC sample of two or more genes selected from those shown in ABT09867-
XX CC ABT1112, where a differential expression of the genes indicates breast
XX CC cancer. The methods are useful in diagnosing, treating, detecting the
XX CC progression, and in monitoring treatment of breast cancer in patients.
XX CC The methods are also useful as a screening tool for agents that modulate
XX CC the onset or progression of breast cancer. The breast cancer genes may be
XX CC used as diagnostic markers for the prediction or identification of the
XX CC malignant state of breast tissue, for confirming the type and progression
XX CC of cancer, and for drug screening and assays. The present sequence is a
XX CC coding sequence of the invention. Note: The sequence data for this patent
XX CC did not form part of the printed specification, but was obtained in
XX CC electronic format directly from WIPO at
XX CC ftp.wipo.int/pub.published_pct_sequences
SQ Sequence 122748 BP; 32088 A; 31056 C; 30547 G; 29057 T; 0 U; 0 Other;
Query Match 38.6%; Score 154.2; DB 6; Length 122748;
Best Local Similarity 75.1%; Pred. No. 2.9e-32;
Matches 208; Conservative 1; Mismatches 59; Indels 9; Gaps 1;
QY 81 ATGCGTGAATCCCGACACTTCGGGAGGCCAAGTGGGCGATCTACCTGAGGTCAAGA 140
DB 114310 ATGCGTGAATCCCGACACTTCGGGAGGCCAAGTGGGCGATCTACCTGAGGTCAAGA 114251
QY 141 TCGAGACCATCTGGCCCAACATGTGAAACCCCGTCTTACTTAAATAACAAAAATAGC 200
DB 114250 TCGAGACCATCTGGCCCAACATGTGAAACCCCGTCTTACTTAAATAACAAAAATAGC 114191
QY 201 TGGGATGATGGGACACACCTCTAGTCCAGCTACTCAGAGCCGGAATTCAGATGAGC 260
DB 114190 TGGGATGATGGGACACCTCTAGTCCAGCTACTCAGAGCCGGAATTCAGATGAGT 114131
QY 261 TGAAGTTCGAGAGTGAAGCGGAATACAGATGACAGAGTGAAGTGAAGTCCGCT 320
DB 114130 TGAAGTTCGAGAGTGAAGCGGAATACAGATGACAGAGTGAAGTGAAGTCCGCT 114080
QY 321 CAAAAACAACAACAAAAACAAAAACCATTAAGACA 357
DB 114079 CAAAAACAACAACAAAAACAAAAACCATTAAGACA 114043
RESULT 13
AAC79009
ID AAC79009 standard; DNA; 884 BP.
XX
AC AAC79009;

XX DT 14-FEB-2001 (first entry)
XX DE Human secreted protein gene 13 clone HSRBCT2.
XX KW Cytostatic; immunosuppressive; neutrotic; neuroprotective; antiviral;
XX KW antiallergic; hepatotropic; antidiabetic; antiinflammatory; antitumor;
XX KW vulnery; anticonvulsant; antibacterial; antifungal; antiparasitic;
XX KW cardiac; gene therapy; cancer; immune disorder; cardiovascular disorder;
XX KW neurological disease; infection; human; secreted protein; ss.
XX OS Homo sapiens.
XX EN WO200058358-A1.
XX PD 05-OCT-2000.
XX PF 23-MAR-2000; 2000WO-US007725.
XX PR 26-MAR-1999; 99US-0126602P.
XX PR 14-JAN-2000; 2000US-0176063P.
XX PA (HUMA-) HUMAN GENOME SCI INC.
XX PI Rosen CA, Ruben SM, Komatsoulis G;
XX DR WPI; 2000-594640/56.
XX DR P-PSDB; AAB44347.
XX PS Fourty nine nucleic acid molecules encoding human secreted proteins,
XX PT useful in the prevention, treatment and diagnosis of cancer, immune
XX PT disorders, cardiovascular disorders and neurological diseases.
XX CC Claim 1; Page 322; 367bp; English.
XX CC The invention relates to the isolation of genes AAC78997-C79045 encoding
XX CC 49 human secreted proteins AAB4435-84482. The genes can be used to
XX CC generate fusion proteins by linking to the gene for the human
XX CC immunoglobulin G Fc portion (AAC78988) for increasing the stability of
XX CC the fusion protein as compared to the human protein only. The genes and
XX CC proteins are useful for preventing, ameliorating or treating medical
XX CC conditions, e.g. by protein or gene therapy. The genes are isolated from
XX CC a range of human tissues disclosed in the specification. The nucleic
XX CC acids, proteins, antibodies and (ant)agonists are useful in the
XX CC diagnosis, treatment and prevention of: (a) cancer, e.g. breast and
XX CC ovarian cancer, and other cancers of the adrenal gland, bone, bone
XX CC marrow, breast, gastrointestinal tract, liver, lung, or urogenital; (b)
XX CC immune disorders e.g. Addison's disease, allergies, autoimmune haemolytic
XX CC anaemia, autoimmune thyroiditis, diabetes mellitus, Crohn's disease,
XX CC multiple sclerosis, rheumatoid arthritis and ulcerative colitis; (c)
XX CC cardiovascular disorders such as myocardial ischaemias; (d) wound healing
XX CC / (e) neurological diseases e.g. cerebral anoxia and epilepsy; and (f)
XX CC infectious diseases such as viral, bacterial, fungal and parasitic
SQ Sequence 884 BP; 211 A; 214 C; 226 G; 233 T; 0 U; 0 Other;
Query Match 38.4%; Score 153.6; DB 3; Length 884;
Best Local Similarity 73.7%; Pred. No. 9.2e-33;
Matches 224; Conservative 1; Mismatches 70; Indels 9; Gaps 2;
QY 37 AATATTAAATGAACATTTGTCGAGCCAGGCACTGCGGATGCTGTAATCCAG 96
DB 585 ATATTATTGTTTAAATGAATCATACAGGCTGGGACAGTGCTCAGCGCTGTAATCTAG 644
QY 97 CACTTCGAGAGCCCAAGGTGGGCGGATCACTGAGGTCAAGAGATGAGACCATCTGAC 156
DB 645 CACTTCGAGAGCCCAAGGTGGGCGGATCACTGAGGTGGGAGTTCGAGACCATCTGAC 704
QY 157 CAACATGTGAACCCCGTCTTTACTTAAATAACAAAAATAGCTGGGCGATGTGGCACA 216
DB 705 CAATATGATGAACACCCCGTCTCTA-AAAAATATACAAAAATTAGCCGGGCGTGTGTGAGG 763

OY		217	CACCTGTATCCAGTACTCAGAGACC	CGGAAATTGCAGAGCTGAAGATTCGAGAGTGA	276
D8		764	CACCTGTATCCAGTACTCAGAGAGCA	GAGGTTTGAGTAGTCAAGTCTGG-----	815
OY		277	GCCGAATCATCAGATCACAGAGTGA	GCAGAGTGAAGAKCKCGTCTCAAAAAACAACAACA	336
D8		816	GGCCATTGCACTCCAGCCTCGGGCA	AAAAAGTGTAAACTCTGTCTCAAAAAAAAAA	875
OY		337	AAAC	340	
D8		876	AAAC	879	
 RESULT 14 AAA64141 ID AAA64141 standard; DNA; 14784 BP. AAA64141 AC AAA64141; XX XX XX DT 20-DEC-2000 (first entry) XX XX D8 Nucleotide sequence of a beta-tubulin antigen. XX XX Beta-tubulin antigen; inner ear protein; Meniere's disease; autoantibody; KV Chronic ear disease; autoimmune disease; ss. XX XX Homo sapiens. OS XX WO2000050593-A1. PN XX 31-AUG-2000. XX XX 25-FEB-2000; 2000WO-US004795. PF XX XX 25-FEB-1999; 99US-0121549P. PR XX PA (UYTE-) UNIV. TENNESSEE RES CORP. XX PI Yoo TU; XX XX WPI; 2000-558400/51. DR XX XX New beta-tubulin antigen in the membranous structure of the inner ear, PT reactive with antibodies of patients with Meniere's disease, for P7 diagnosing Meniere's disease and distinguishing this disease from other P1 autoimmune ear diseases. XX XX XX Claim 3; Page 97-103; 115pp; English. PS XX XX The present sequence encodes a beta-tubulin antigen. The protein is an CC antigen of the membranous structure of the inner ear protein, and is CC reactive with antibodies from patients having Meniere's disease. CC Meniere's disease is a chronic ear disease with unknown etiology. Serum CC from patients suffering from this disease contain autoantibodies against CC a 30 kDa cochlear protein antigen. The disease is believed to be an CC autoimmune disease. The beta-tubulin antigen is useful as a target CC substance in diagnosing or detecting Meniere's disease and in CC distinguishing this disease from other autoimmune ear diseases CC SO Sequence 14784 BP; 5454 A; 2966 C; 2926 G; 3438 T; 0 U; 0 Other;					
OY		Query Match	38.3%; Score 153.2; DB 3; Length 14784;		
D8		Best Local Similarity	65.1%; Pred. No. 2.9e-32;		
		Matches	241; Conservative 0; Mismatches 128; Indels 1; Gaps 1		
OY		31	GAACCAAATAATTAATAAGACATTGT	CAGGCCAGGCATGACACTGGCTGAATCCTGTAA	90
D8		3372	GATCAAGAAATTTATTTATTAAGTGA	GCAAGCGCTGGCGCGGTCCACGCGCTGTAA	3433
OY		91	TCCCAAGCACTTCGGAGAGCCAAAGT	GGCGGATCACTTGAGTCAAGAAGATGAAGACCT	150
D8		3432	TCCCAAGCACTTCGGAGAGCCAAAGT	GGCGGATCACTTGAGTCAAGAAGATGAAGACCA	3493

Qy	151	CCTGGCGCAACATGTGTGAAACCCCGCTTTACTATAAAATACAAAAATAGTGGCATGCT	21
Db	3492	CTTGCGCAACATGTGGCAAAACCCCGCTTCTACTAAAAATACAAAAATTAGCGACATGCT	35
Qy	211	GGCACACACCTGTAGTCCAGCTACTCAGAGCCGAGATTGCAGTGAGTGAATGCA	270
Db	3552	GGCGCACACCTGTAGTCCAGCTACTCAGAGGCTTAGGACAGAGAACCACTGAACCCG	361
Qy	271	GAGTAGCCGAATATCACAGATCACAGAGTGAGCAGATGAGACCCGTCCTCAAAACAC	330
Db	3612	GGAGGTGTGTGATGACAGTGAGCCGAGATGTGTG-CATTGCAACGCCAGCCTGCACAACAG	367
Qy	331	AACAAAAAACAAAAAACCATATAAGCATGTCTCATCTGCGGCTCCAGACTATATTGAGGA	390
Db	3671	ACCAAAAAAACAAAAAATTGTGGCCAGTACAAAACTTCATTATAATACATTGCAATTT	3737
Qy	391	GACCAAAAAG 400	
Db	3731	TTCCATCCAG 3740	
RESULT 15			
AE03362			
ID	AE03362	strand; DNA; 166942 BP.	
XX			
AC	AE03362;		
XX			
DT	09-FEB-2006 (first entry)		
XX			
DE	Acute myelogenous leukemia prognosis related DNA sequence SEQ ID NO: 23.		
XX			
KM	ds; gene; acute myelogenous leukemia; prognosis; gene expression;		
XX	biochip.		
XX			
OS	Homo sapiens.		
XX			
PN	JP200533987-A.		
PD			
PD	08-DEC-2005.		
XX			
PF	06-MAY-2005; 2005JP-00135284.		
XX			
PR	06-MAY-2004; 2004US-0568635P.		
XX			
PA	(VERI-) VERIDEX LLC.		
XX			
PI	Laponi M;		
XX			
DR	WPI, 2006-004067/01.		
XX			
PT			
PT	Evaluating an acute myelogenous leukemia patient's prognosis, comprising		
PT	detecting lower or higher expression level of gene recognized by probe		
PT	set e.g. 202820-at and 206148-at, with respect to predetermined cut-off		
XX	level.		
XX			
PS	Example 5; SEQ ID NO 23; 60bp; Japanese.		
XX			
CC	The present sequence is that of a human gene sequence which is claimed		
CC	for use in evaluating the prognosis of patients suffering from acute		
CC	myelogenous leukemia (AML) by analysis of the patients gene expression		
CC	profile. The invention relates to a novel method for evaluating the		
CC	prognosis of a patient with acute myelogenous leukemia by detecting		
CC	higher/lower expression level of genes encoding mRNA recognized by a		
CC	probe set chosen from 19 nucleotide sequences (AE03340 or AE03344-		
CC	AE03361). Also, gene expression profiling of AML patients using a set of		
CC	probes based on 167 sequences (AE03362-AE03527) using a biochip was		
CC	performed to determine prognosis. Methods are also included for		
CC	determining an AML patient's state, determining AML patient's treatment		
CC	protocol, determining whether the patient will respond to the treatment,		
CC	and producing an AML patient's prognosis report by analysis of the		
CC	patients gene expression profile. The methods are useful in evaluating an		
CC	AML patient's prognosis and for treating AML patient.		

[illegible]

PR 08-NOV-2000; 2000US-0246528P.
PR 08-NOV-2000; 2000US-0246532P.
PR 08-NOV-2000; 2000US-0246609P.
PR 08-NOV-2000; 2000US-0246610P.
PR 08-NOV-2000; 2000US-0246611P.
PR 08-NOV-2000; 2000US-0246613P.
PR 17-NOV-2000; 2000US-0249207P.
PR 17-NOV-2000; 2000US-0249208P.
PR 17-NOV-2000; 2000US-0249209P.
PR 17-NOV-2000; 2000US-0249210P.
PR 17-NOV-2000; 2000US-0249211P.
PR 17-NOV-2000; 2000US-0249212P.
PR 17-NOV-2000; 2000US-0249213P.
PR 17-NOV-2000; 2000US-0249214P.
PR 17-NOV-2000; 2000US-0249215P.
PR 17-NOV-2000; 2000US-0249216P.
PR 17-NOV-2000; 2000US-0249217P.
PR 17-NOV-2000; 2000US-0249218P.
PR 17-NOV-2000; 2000US-0249244P.
PR 17-NOV-2000; 2000US-0249245P.
PR 17-NOV-2000; 2000US-0249264P.
PR 17-NOV-2000; 2000US-0249265P.
PR 17-NOV-2000; 2000US-0249297P.
PR 17-NOV-2000; 2000US-0249300P.
PR 01-DEC-2000; 2000US-0250160P.
PR 01-DEC-2000; 2000US-0250391P.
PR 05-DEC-2000; 2000US-0251030P.
PR 05-DEC-2000; 2000US-0251030P.
PR 05-DEC-2000; 2000US-0251888P.
PR 06-DEC-2000; 2000US-0251479P.
PR 08-DEC-2000; 2000US-0251856P.
PR 08-DEC-2000; 2000US-0251868P.
PR 08-DEC-2000; 2000US-0251869P.
PR 08-DEC-2000; 2000US-0251989P.
PR 08-DEC-2000; 2000US-0251990P.
PR 11-DEC-2000; 2000US-0254097P.
PR 05-JAN-2001; 2001US-0259678P.
XX
XX (HUMA-) HUMAN GENOME SCI INC.
XX
XX Rosen CA, Barash SC, Ruben SM;
XX WPI; 2001-465570/50.
XX
XX Isolated nucleic acid molecule encoding a reproductive system antigen is
XX used in preventing, treating or ameliorating a medical condition.
XX
XX Disclosure; SEQ ID NO 7647; 1297pp + Sequence Listing; English.
XX
XX The present invention provides the protein and coding sequences of a
XX number of human reproductive system related antigens. These can be used
XX in the prevention and treatment of reproductive system disorders,
XX including cancer. The present sequence is a genomic sequence encoding a
XX protein of the invention
XX
SQ Sequence 4388 BP; 870 A; 1238 C; 1205 G; 1075 T; 0 U; 0 Other;
Query Match 38.1%; Score 152.4; DB 4; Length 4388;
Best Local Similarity 76.5%; Pred. No. 3.3e-32;
Matches 202; Conservative 1; Mismatches 52; Indels 9; Gaps 1;
OY 81 ATGCTGTAATCCGACGACTTCGGAGGCCGAGGCGGATCCTGAGGTCAAGAGA 140
DB 1275 AAGCTGTAATCCGACGACTTCGGAGGCCGAGGCGGATCCTGAGGTCAAGAGA 1216
OY 141 TCGAGACATCTGGCGCAACATGAGAAACCCGCTTACTTAATAAAATACAAAATATGC 200
DB 1215 TCGAGACATCTGGCGCAACATGAGAAACCCGCTTACTTAATAAAATACAAAATATGC 1156
OY 201 TGGGATGATGGGACACACCTGTAGTCCAGACTACTCAGAGCGGAGATTGCAGTGAGC 260
DB 1155 CGCACGTGTGGCGGCGGCTGTATCCCACTACTCGGAGGAGGAGGTTCAGCGAGC 1096

OY 261 TGAATCCGACAGTGAAGCCGAATTCACATCAGAGTGAACGACGACCCGCTT 320
DB 1095 AGAGATCGTCCGCTTGCACCTCCAGTCTGGGCGA-----CACAGGAGACTCCGCTT 1045
OY 321 CAAATACACACACAAATACAAATACAA 344
DB 1044 CAAATACACACACAAATACAAATACAA 1021
RESULT 17
ABL97853/c
ID ABL97853 standard; DNA; 4388 BP.
XX
XX ABL97853;
XX
XX 21-JUN-2002 (first entry)
XX
XX Human testicular antigen encoding DNA fragment SEQ ID NO: 2505.
XX
XX Human; testicular antigen; testes; cancer; metastasis; immune disorder;
XX reproductive system disorder; urinary system disorder; gene therapy;
XX cardiovascular disorder; respiratory disorder; neurological disorder;
XX gastrointestinal disease; infection; cytostatic; gene; ds.
OS Homo sapiens.
XX
XX WO200155317-A2.
XX
XX 02-AUG-2001.
XX
XX 17-JAN-2001; 2001WO-US001329.
XX
XX 31-JAN-2000; 2000US-0179065P.
XX 04-FEB-2000; 2000US-0180628P.
XX 24-FEB-2000; 2000US-0184664P.
XX 02-MAR-2000; 2000US-0186350P.
XX 16-MAR-2000; 2000US-0189874P.
XX 17-MAR-2000; 2000US-0190076P.
XX 18-APR-2000; 2000US-0198123P.
XX 19-MAY-2000; 2000US-0205515P.
XX 07-JUN-2000; 2000US-0209467P.
XX 28-JUN-2000; 2000US-0214886P.
XX 30-JUN-2000; 2000US-0215135P.
XX 07-JUL-2000; 2000US-0216647P.
XX 11-JUL-2000; 2000US-0216880P.
XX 11-JUL-2000; 2000US-0217487P.
XX 14-JUL-2000; 2000US-0217496P.
XX 14-JUL-2000; 2000US-0218290P.
XX 26-JUL-2000; 2000US-0220963P.
XX 26-JUL-2000; 2000US-0220964P.
XX 14-AUG-2000; 2000US-0224518P.
XX 14-AUG-2000; 2000US-0224519P.
XX 14-AUG-2000; 2000US-0225213P.
XX 14-AUG-2000; 2000US-0225214P.
XX 14-AUG-2000; 2000US-0225214P.
XX 14-AUG-2000; 2000US-0225266P.
XX 14-AUG-2000; 2000US-0225267P.
XX 14-AUG-2000; 2000US-0225268P.
XX 14-AUG-2000; 2000US-0225270P.
XX 14-AUG-2000; 2000US-0225447P.
XX 14-AUG-2000; 2000US-0225757P.
XX 14-AUG-2000; 2000US-0225757P.
XX 14-AUG-2000; 2000US-0225758P.
XX 14-AUG-2000; 2000US-0225759P.
XX 18-AUG-2000; 2000US-0226279P.
XX 22-AUG-2000; 2000US-0226681P.
XX 22-AUG-2000; 2000US-0226686P.
XX 22-AUG-2000; 2000US-0227182P.
XX 23-AUG-2000; 2000US-0227009P.
XX 30-AUG-2000; 2000US-0228924P.
XX 01-SEP-2000; 2000US-0229287P.
XX 01-SEP-2000; 2000US-0229343P.
XX 01-SEP-2000; 2000US-0229344P.
XX 01-SEP-2000; 2000US-0229345P.

Query Match	38.1%; Score 152.4; DB 4; Length 4388;
Best Local Similarity	76.5%; Pred. No. 3.3e-32;
Matches 202; Conservative 1; Mismatches 52; Indels 9; Gaps 1;	
QY 81 ATGCGTGAATCCCAACACTTCGGGAGGCCAAGTGGGGCGATCCTGAGCTCAAGAGA 140	
DB 1275 AAGCCTGTAATTCAGACACTTTGGGAGGCCGAGGCCGCGGATCACTTGAGGTGAGGAGT 1216	
QY 141 TCGAGACCATCTGGCCCAACATGGTGAACCCCGTCTTTACTAAATAACAAAATATAGC 200	
DB 1215 TCGAGACCATCTGGCCCAACATGGTGAACCCCGTCTTTACTAAATAACAAAATATAGC 1156	
QY 201 TGGGATGATGGGACACACACTGTAGTCCAGACTACTGAGAGCCGAGATTGCAGTAGC 260	
DB 1155 CGCAGTGTGTGGCGCGCCCTGTATATCCAGCTACTCGGGAGGCGGAGGTTGACAGCGAGC 1096	
QY 261 TGAAGTCGACAGTGAAGCCGAATCAACAGATCACAGAGTGAAGAGTGAAGCCGCTCT 320	
DB 1095 AGAGATCGTCGATTCAGCTCACCTCACTCTGGGCGA-----CACAGCGAGACTCGCTCT 1045	
QY 321 CAAAACACACACACAAAACAAA 344	
DB 1044 CAAAAAAAAAAAAAAAAAAAAA 1021	

RESULT 18
 ABL68122/c
 ID ABL68122 standard; DNA; 174424 BP.

XX AB68122;
 XX
 DT 15-MAY-2002 (first entry)
 XX
 DE Ovary cancer related gene sequence SEQ ID NO:6459.
 XX
 KY Human; cancer: colon; breast; ovary; oesophagus; kidney; thyroid;
 KY stomach; lung; prostate; pancreas; carcinoma; antitumour; cancerous;
 KY cytostatic; gene therapy; antineoplastic; Wilms tumour; adenocarcinoma;
 KY gene; ds.
 XX
 OS Homo sapiens.
 XX
 PN WO200194629-A2.
 XX
 PD 13-DEC-2001.
 XX
 PE 30-MAY-2001; 2001WO-US010838.
 XX
 PR 05-JUN-2000; 2000US-0209473P.
 PR 05-JUN-2000; 2000US-0209531P.
 PR 18-SEP-2000; 2000US-0231133P.
 PR 18-SEP-2000; 2000US-0231617P.
 PR 20-SEP-2000; 2000US-0234009P.
 PR 20-SEP-2000; 2000US-0234034P.
 PR 20-SEP-2000; 2000US-0234052P.
 PR 22-SEP-2000; 2000US-0234509P.
 PR 22-SEP-2000; 2000US-0234567P.
 PR 25-SEP-2000; 2000US-0234923P.
 PR 25-SEP-2000; 2000US-0234924P.
 PR 25-SEP-2000; 2000US-0235077P.
 PR 25-SEP-2000; 2000US-0235082P.
 PR 25-SEP-2000; 2000US-0235134P.
 PR 25-SEP-2000; 2000US-0235280P.
 PR 26-SEP-2000; 2000US-0235637P.
 PR 26-SEP-2000; 2000US-0235638P.
 PR 27-SEP-2000; 2000US-0235711P.
 PR 27-SEP-2000; 2000US-0235720P.
 PR 27-SEP-2000; 2000US-0235840P.
 PR 27-SEP-2000; 2000US-0235863P.
 PR 28-SEP-2000; 2000US-0236028P.
 PR 28-SEP-2000; 2000US-0236032P.
 PR 28-SEP-2000; 2000US-0236033P.
 PR 28-SEP-2000; 2000US-0236034P.
 PR 28-SEP-2000; 2000US-0236109P.
 PR 28-SEP-2000; 2000US-0236111P.
 PR 29-SEP-2000; 2000US-0236842P.
 PR 29-SEP-2000; 2000US-0236891P.
 PR 02-OCT-2000; 2000US-0237172P.
 PR 02-OCT-2000; 2000US-0237173P.
 PR 02-OCT-2000; 2000US-0237278P.
 PR 02-OCT-2000; 2000US-0237294P.
 PR 02-OCT-2000; 2000US-0237295P.
 PR 02-OCT-2000; 2000US-0237316P.
 PR 03-OCT-2000; 2000US-0237425P.
 PR 03-OCT-2000; 2000US-0237598P.
 PR 03-OCT-2000; 2000US-0237604P.
 PR 03-OCT-2000; 2000US-0237606P.
 PR 03-OCT-2000; 2000US-0237608P.
 PR 01-NOV-2000; 2000US-0244867P.
 PR 01-NOV-2000; 2000US-0245084P.
 XX
 PA (AVAL-) AVALON PHARM.
 XX
 PI Young PE, Augustus M, Carter KC, Ebner R, Endress G, Horrigan S;
 PI Sopet DR, Weaver Z;
 XX
 DR WPI; 2002-188264/24.
 XX
 PT Screening for anti-neoplastic agent involves exposing cells to a chemical
 PT agent to be tested for anti-neoplastic activity, and determining a change
 PT in expression of a gene of a signature gene set.

XX Claim 1; SEQ ID NO 6459; 44bp; English.
 PS
 CC The present invention describes a method (M1) for screening for an anti-
 CC neoplastic agent. The method involves exposing cells to a chemical agent
 CC to be tested for anti-neoplastic activity, determining a change in
 CC expression of at least one gene (I) of a signature gene set, where (I)
 CC comprises a sequence (S) selected from 8447 sequences (given in AB61664
 CC to AB170110), or is at least 95% identical to (S), where a change in
 CC expression is indicative of anti-neoplastic activity. (I) has cytostatic
 CC activity and can be used in gene therapy. M1 can be used for screening an
 CC anti-neoplastic agent, and can be used for producing a product which is
 CC the data collected with respect to the anti-neoplastic agent as a result
 CC of M1, and the data is sufficient to convey the chemical structure and/or
 CC properties of the agent. M1 can be used in the treatment of cancer such
 CC as colon, breast, stomach, lung, thyroid, oesophageal, ovarian, kidney,
 CC prostate or pancreatic cancer, adenocarcinoma, carcinoma, clear cell
 CC cancer, infiltrating ductal cancer, infiltrating lobular cancer, squamous
 CC cell carcinoma, neuroendocrine carcinoma, papillary carcinoma and Wilms
 CC tumour.
 CC
 SQ Sequence 174424 BP; 39582 A; 48304 C; 48535 G; 38003 T; 0 U; 0 Other;
 XX
 XX
 XX Query Match 38.0%; Score 152; DB 6; Length 174424;
 XX Best Local Similarity 71.8%; Pred. No. 1.3e-31;
 XX Matches 214; Conservative 0; Mismatches 80; Indels 4; Gaps 1;
 XX
 QY 6 TACTCAGCCATGTCCTGGCCATGSGAACCATAATATTAATAGACATTGTCAGGCCAG 65
 DB 48438 TCCTCAGGTGTCTGTATCTTGTGTCTTCTCATTTTAAGATGGGCGAGCTGG 48379
 QY 66 CATGACACTGGCTGAATCCTGTATATCCAGCACTTCGGAGGCCAAGTGGCGGATCA 125
 DB 48378 CATGCA---GCTCATGCTCTGTATCCCAACTTGGAGGCGAAGTGGCGGATCA 48323
 QY 126 CCTGAGGTCAGAGATGAGACATCTCTGGCAACATGTAACCCGCTCTTAATAA 185
 DB 48322 CTTGAGGTCAGAGATCAAGTCATCTCTGGCAACATGTAACCCGCTCTTAATAA 48263
 QY 186 AATACAAAAATAGCTGGGATGTTGGCACACACTGTAGTCCAGCTACTCAGGAGCG 245
 DB 48262 AATACAAAAATAGCTGGGATGTTGGCACACACTGTAGTCCAGCTACTCAGGAGCG 48203
 QY 246 GAGATTGCACTGACCTGATGATCGAGGTGACCGGAATCAAGATCAAGAGTGAGC 303
 DB 48202 GAGCAGGAGAAATCGTTGAATCCGGAGGTGGAGATTGCAGTGAGCCGAGATTGTGC 48145
 XX
 XX RESULT 19
 XX ADQ19573/c
 XX ID ADQ19573 standard; DNA; 181343 BP.
 XX
 XX AC ADQ19573;
 XX
 XX DT 26-AUG-2004 (first entry)
 XX
 XX DE Human soft tissue sarcoma-upregulated DNA - SEQ ID 2392.
 XX
 XX DE soft tissue sarcoma; cytostatic; gene therapy; vaccine; screening; human;
 XX ds.
 XX
 XX OS Homo sapiens.
 XX
 XX PN WO2004048938-A2.
 XX
 XX PD 10-JUN-2004.
 XX
 XX PE 26-NOV-2003; 2003WO-US038193.
 XX
 XX PR 26-NOV-2002; 2002US-0429739P.
 XX
 XX PA (PROT-) PROTEIN DESIGN LABS INC.
 XX

PI Aziz N, Gineburg MM, Zlotnik A;

XX WPI; 2004-441208/41.

PT Early detection of soft tissue sarcoma comprises determining expression
PT of a gene in a first soft tissue sample and a normal soft tissue sample
PT and comparing the gene expression, also useful in treating soft tissue
PT sarcoma.

XX Example 2; SEQ ID NO 2392; 210bp; English.

XX The invention relates to a novel method for detecting soft tissue sarcoma
XX which comprises obtaining a first soft tissue sample from an individual
XX and a normal soft tissue sample from the same or different individual,
XX determining the expression of a gene in both samples and comparing the
XX expression of the gene in both soft tissue samples, where a higher level
XX of protein expression in the first soft tissue sample indicates the
XX presence of soft tissue sarcoma. The method of the invention has
XX cyrostatic applications and may be useful for detecting soft tissue
XX sarcoma, possibly via gene therapy or vaccine production. The nucleic
XX acid sequences may be useful in diagnostic and screening applications.
XX The current sequence is that of a human soft tissue sarcoma-upregulated
XX DNA of the invention. The current sequence is not shown within the
XX specification per se but was submitted in CD format by the inventor.

XX Sequence 181343 BP; 41465 A; 49982 C; 50309 G; 39587 T; 0 U; 0 Other;

XX Query Match 38.0%; Score 152; DB 12; Length 181343;

XX Best Local Similarity 71.8%; Pred. No. 1.3e-31;

XX Matches 214; Conservative 0; Mismatches 80; Indels 4; Gaps 1;

XX 6 TATCTACGACATGCTGTGGCCATGGAGCCCAATATTAATGAACATTCACGGCCG 65

XX 48439 TCTCAGGTGTGCTGTATCTTGTCTTCTCAATTTAAGATGGGGAGGCTGGG 48380

XX 66 CATGACACTGCTGATCTGTATCCAGACACTTCGGAGGCCAAGTGGGCGATCA 125

XX 48379 CATGCA----GCTCATGCTGTATCCCAACACTTTGGAGGCCGAGTGGGATCA 48324

XX 126 CTTGAGGTCAAGATCGAGACCATCTCTGGCCACATGCTGTAATCTTAA 185

XX 48323 CCTGAGGTCAAGATCGAGACCATCTCTGGCCACATGCTGTAATCTTAA 48264

XX 186 AATCAAAATATGCTGGGATGTCGACACCTGTAGTCCGACTACAGGAGCG 245

XX 48263 AATCAAAATATGCTGGGATGTCGACACCTGTAGTCCGACTACAGGAGCG 48204

XX 246 GAGATTGCTGAGCTGAGATCGACAGTGAAGCCGAAATCAAGATCAAGAGTGA 303

XX 48203 GAGGAGAGAAATCGCTTGAATCCGGAGGTGAGATTGCAAGTGAAGCCGAGATTGTGC 48146

RESULT 20

AB08186/c ID AB08186 standard; cDNA; 169739 BP.

XX AB08186;

XX 18-SEP-2002 (first entry)

XX Human osteoblast differentiation related cDNA SEQ ID NO 93.

XX Human, osteoblast; stem cell differentiation; bone tissue deposition;

XX osteoporosis; osteopathic; ss.

XX Homo sapiens.

XX WO200250301-A2.

XX 27-JUN-2002.

XX 18-DEC-2001; 2001WO-US048276.

PR 18-DEC-2000; 2000US-0255882P.

PR 24-APR-2001; 2001US-0285691P.

XX (GENE-) GENE LOGIC INC.

XX (PROC) PROCTER & GAMBLE CO.

XX J. D. Axelrod DW, Cook JS, Jaiswal N, Einstein R, Houghton A;

XX WPI; 2002-557663/59.

XX Use of genes and their expression profiles associated with osteoblast
XX differentiation for screening modulators bone formation, for diagnosing
XX or treating e.g. osteoporosis, or as markers for the differentiation
XX process.

XX Claim 1; SEQ ID NO 93; 78bp + Sequence Listing; English.

XX The invention relates to genes and their expression profiles are used
XX for: (a) screening modulators of precursor stem cell differentiation into
XX osteoblasts, or bone tissue deposition; (b) diagnosing abnormal
XX deposition of bone tissue, abnormal rate of osteoblast formation or
XX osteoporosis; or (c) treating or monitoring treatment of the conditions
XX cited in (b), or monitoring the progression of bone tissue deposition.
XX Specific conditions include postmenopausal osteoporosis, glucocorticoid
XX osteoporosis or male osteoporosis, osteopenia, osteodystrophy, drug-
XX induced abnormalities in bone formation or bone loss, conditions that
XX involve altered bone metabolism (e.g. idiopathic juvenile osteoporosis),
XX skeletal disease linked to breast cancer, mastocytosis, Fanconi syndrome
XX or fibrous dysplasia. The present sequence is that of an osteoblast
XX differentiation associated cDNA marker of the invention. Note: The
XX sequence data for this patent did not form part of the printed
XX specification, but was obtained in electronic format directly from WIPO
XX at ftp.wipo.int/pub/published_pct_sequences

XX Sequence 169739 BP; 49809 A; 35660 C; 35715 G; 48555 T; 0 U; 0 Other;

XX Query Match 37.9%; Score 151.4; DB 6; Length 169739;

XX Best Local Similarity 70.9%; Pred. No. 1.9e-31;

XX Matches 229; Conservative 1; Mismatches 87; Indels 6; Gaps 2;

XX 30 GAATCCAAATATTAATGAACATTCAGGCCAGGACATGCTGCTGAATCTGTGA 89

XX 125892 GGTAAAGAAATCTTAAGTAAAGATCTAAAGTTTGGCCAGGACATGAGCTGAACCTGTGA 125833

XX 90 ATCCAGACATCTGGGAGGCCCAAGTGGCGGATCCTGAGTCAGAGATCGAGACCA 149

XX 125832 ATCCAGACATCTGGGAGGCCCAAGTGGCGGATCCTGAGTCAGAGATCGAGACCA 125773

XX 150 TCTGGCCAAATGATGTAATCCCGCTCTTAAATCAAAAAATAGCTGGGCAATG 209

XX 125772 GCTGGCCAAATGATGTAATCCCGCTCTTAAATCAAAAAATAGCTGGGCAATG 125713

XX 210 TGGACACACCTGTAGTCCAGTACTCAGAGCCGAGATGTCAGTGAAGATGCTGC 269

XX 125712 TGGACACACCTGTAGTCCAGTACTCAGAGCCGAGATGTCAGTGAAGATGCTGC 125654

XX 270 AGATGAGCCGAATCA-----GATCAAGATGAGCAGAGTGAAGCCKCGTCTCAAA 324

XX 125653 TGGGTGATCAAGATCGACATGCGCACTCCAGCTTGGGCAACGAGATCTCATTCGAA 125594

XX 325 AACCAACCAAAAAA 347

XX 125593 AAAAATTAATTAATTAATTAATTA 125571

RESULT 21

AAK68418/c ID AAK68418 standard; DNA; 14282 BP.

XX AAK68418;

XX 06-NOV-2001 (first entry)

XX Human immune/haematopoietic antigen genomic sequence SEQ ID NO:23230.
XX Human; immune; haematopoietic; immune/haematopoietic antigen; cancer;
XX Cytostatic; gene therapy; vaccine; metastasis; ds.
XX Homo sapiens.
XX MO200157182-A2.
XX 09-AUG-2001.
XX 17-JAN-2001; 2001WO-US001354.
XX 31-JAN-2000; 2000US-0179065P.
XX 04-FEB-2000; 2000US-0180628P.
XX 24-FEB-2000; 2000US-0184664P.
XX 02-MAR-2000; 2000US-0186350P.
XX 16-MAR-2000; 2000US-0189874P.
XX 17-MAR-2000; 2000US-0190076P.
XX 18-APR-2000; 2000US-0198123P.
XX 19-MAY-2000; 2000US-0205515P.
XX 07-JUN-2000; 2000US-0209467P.
XX 28-JUN-2000; 2000US-0215135P.
XX 30-JUN-2000; 2000US-0216647P.
XX 07-JUL-2000; 2000US-0216880P.
XX 11-JUL-2000; 2000US-0217487P.
XX 11-JUL-2000; 2000US-0217496P.
XX 14-JUL-2000; 2000US-0218290P.
XX 26-JUL-2000; 2000US-0220963P.
XX 14-AUG-2000; 2000US-0224518P.
XX 14-AUG-2000; 2000US-0225267P.
XX 14-AUG-2000; 2000US-0225268P.
XX 14-AUG-2000; 2000US-0225270P.
XX 14-AUG-2000; 2000US-0225447P.
XX 14-AUG-2000; 2000US-0225757P.
XX 14-AUG-2000; 2000US-0225758P.
XX 14-AUG-2000; 2000US-0225759P.
XX 18-AUG-2000; 2000US-0226279P.
XX 22-AUG-2000; 2000US-0226681P.
XX 22-AUG-2000; 2000US-0226868P.
XX 23-AUG-2000; 2000US-0227182P.
XX 30-AUG-2000; 2000US-0228924P.
XX 01-SEP-2000; 2000US-0229287P.
XX 01-SEP-2000; 2000US-0229343P.
XX 01-SEP-2000; 2000US-0229345P.
XX 05-SEP-2000; 2000US-0229509P.
XX 05-SEP-2000; 2000US-0229513P.
XX 06-SEP-2000; 2000US-0230437P.
XX 06-SEP-2000; 2000US-0230438P.
XX 08-SEP-2000; 2000US-0231243P.
XX 08-SEP-2000; 2000US-0231244P.
XX 08-SEP-2000; 2000US-0231244P.
XX 08-SEP-2000; 2000US-0231413P.
XX 08-SEP-2000; 2000US-0231414P.
XX 08-SEP-2000; 2000US-0232080P.
XX 12-SEP-2000; 2000US-0232081P.
XX 14-SEP-2000; 2000US-0232397P.
XX 14-SEP-2000; 2000US-0232398P.
XX 14-SEP-2000; 2000US-0232399P.
XX 14-SEP-2000; 2000US-0232400P.
XX 14-SEP-2000; 2000US-0232401P.
XX 14-SEP-2000; 2000US-0233063P.
XX 14-SEP-2000; 2000US-0233064P.
PR 14-SEP-2000; 2000US-0233065P.
PR 21-SEP-2000; 2000US-0234223P.
PR 21-SEP-2000; 2000US-0234274P.
PR 25-SEP-2000; 2000US-0234979P.
PR 25-SEP-2000; 2000US-0234980P.
PR 26-SEP-2000; 2000US-0235484P.
PR 27-SEP-2000; 2000US-0235834P.
PR 27-SEP-2000; 2000US-0235836P.
PR 29-SEP-2000; 2000US-0236327P.
PR 29-SEP-2000; 2000US-0236367P.
PR 29-SEP-2000; 2000US-0236368P.
PR 29-SEP-2000; 2000US-0236369P.
PR 29-SEP-2000; 2000US-0236370P.
PR 02-OCT-2000; 2000US-0236802P.
PR 02-OCT-2000; 2000US-0237037P.
PR 02-OCT-2000; 2000US-0237038P.
PR 02-OCT-2000; 2000US-0237039P.
PR 02-OCT-2000; 2000US-0237040P.
PR 13-OCT-2000; 2000US-0239935P.
PR 13-OCT-2000; 2000US-0239937P.
PR 20-OCT-2000; 2000US-0240960P.
PR 20-OCT-2000; 2000US-0241221P.
PR 20-OCT-2000; 2000US-0241785P.
PR 20-OCT-2000; 2000US-0241787P.
PR 20-OCT-2000; 2000US-0241808P.
PR 20-OCT-2000; 2000US-0241809P.
PR 20-OCT-2000; 2000US-0241826P.
PR 01-NOV-2000; 2000US-0244617P.
PR 08-NOV-2000; 2000US-0246474P.
PR 08-NOV-2000; 2000US-0246475P.
PR 08-NOV-2000; 2000US-0246476P.
PR 08-NOV-2000; 2000US-0246477P.
PR 08-NOV-2000; 2000US-0246478P.
PR 08-NOV-2000; 2000US-0246523P.
PR 08-NOV-2000; 2000US-0246524P.
PR 08-NOV-2000; 2000US-0246525P.
PR 08-NOV-2000; 2000US-0246526P.
PR 08-NOV-2000; 2000US-0246527P.
PR 08-NOV-2000; 2000US-0246528P.
PR 08-NOV-2000; 2000US-0246532P.
PR 08-NOV-2000; 2000US-0246609P.
PR 08-NOV-2000; 2000US-0246610P.
PR 08-NOV-2000; 2000US-0246611P.
PR 08-NOV-2000; 2000US-0246613P.
PR 17-NOV-2000; 2000US-0249207P.
PR 17-NOV-2000; 2000US-0249208P.
PR 17-NOV-2000; 2000US-0249209P.
PR 17-NOV-2000; 2000US-0249210P.
PR 17-NOV-2000; 2000US-0249211P.
PR 17-NOV-2000; 2000US-0249212P.
PR 17-NOV-2000; 2000US-0249213P.
PR 17-NOV-2000; 2000US-0249214P.
PR 17-NOV-2000; 2000US-0249215P.
PR 17-NOV-2000; 2000US-0249216P.
PR 17-NOV-2000; 2000US-0249217P.
PR 17-NOV-2000; 2000US-0249218P.
PR 17-NOV-2000; 2000US-0249219P.
PR 17-NOV-2000; 2000US-0249300P.
PR 01-DEC-2000; 2000US-0250160P.
PR 01-DEC-2000; 2000US-0250391P.
PR 05-DEC-2000; 2000US-0251030P.
PR 05-DEC-2000; 2000US-0251988P.
PR 06-DEC-2000; 2000US-0251479P.
PR 08-DEC-2000; 2000US-0251856P.
PR 08-DEC-2000; 2000US-0251868P.
PR 08-DEC-2000; 2000US-0251869P.

PR 08-DEC-2000; 2000US-0251989P.
PR 08-DEC-2000; 2000US-0251990P.
PR 11-DEC-2000; 2000US-0254057P.
PR 05-JAN-2001; 2001US-0259678P.
XX
XX (HUMA-) HUMAN GENOME SCI INC.
PI Rosen CA, Barash SC, Ruben SM;
DR WPI; 2001-483426/52.
XX
XX Nucleic acids encoding human immune/hematopoietic antigen polypeptides,
PT useful for preventing, diagnosing and/or treating cancers and metastasis.
XX
XX Disclosure; SEQ ID NO 23230; 3071bp + Sequence Listing; English.
PS
XX AAK54951 to AAK64702 encode the human immune/haematopoietic antigen (I)
CC amino acid sequences given in AAM82170 to AAM91921. (I) have cytostatic
CC activity, and can be used in gene therapy and vaccine production. (I)
CC proteins and polynucleotides may be used in the prevention, diagnosis and
CC treatment of diseases associated with inappropriate (I) expression. For
CC example, they may be used to treat disorders associated with decreased
CC expression by rectifying mutations or deletions in a patient's genome
CC that affect the activity of (I) by expressing inactive proteins or to
CC supplement the patients own production of (I). Additionally, (I)
CC polynucleotides may be used to produce the secreted (I), by inserting the
CC nucleic acids into a host cell and culturing the cell to express the
CC protein. (I) proteins and polynucleotides may be used to prevent,
CC diagnose and treat immune/haematopoietic-related diseases, especially
CC cancers and cancer metastases of haematopoietic-derived cells. AAK64703
CC to AAK87694 represent human immune/haematopoietic antigen genomic
CC sequences from the present invention. AAK54942 to AAK54950 and AAM82169
CC represent sequences used in the exemplification of the present invention
XX
XX Sequence 14282 BP; 3993 A; 2449 C; 2814 G; 5026 T; 0 U; 0 Other;
SQ
Query Match 37.7%; Score 150.6; DB 4; Length 14282;
Best Local Similarity 73.6%; Pred. No. 1.5e-31;
Matches 192; Conservative 0; Mismatches 65; Indels 0; Gaps 0;
QY 48 AGACATTGTCAGGCGGAGCATGACACTGGCTGAATGCTGTAATCCGACACTTCGGAG 107
DB 4784 AATATATGTATCTCTGCGGACGACGCGGGGCAATGCTGTAATCCGACACTTCGGAG 4725
QY 108 GCCAAGTGGGCGGATGATCCTGAGGTCAAGAGATGAGACCATCTGGCCAAATGTGGA 167
DB 4724 ACCGAGTGGGCGGATGATCCTGAGGTGAGAGTTCAGGCTGCGGCAATGTGGA 4665
QY 168 AACCCCGCTTAACTAAATAATACAAAATAATAGCTGGGATGATGACACACTGTAGTC 227
DB 4664 AACCCCGCTTAACTAAATAATAGCTGGGATGATGACACACTGTAGTC 4605
QY 228 CCAGCTACTCAGAGCGCGAGATTGAGTGAAGCTGAGATTCGACAGTGAAGCCGAATCAC 287
DB 4604 CCAGCTACTCAGAGCGCGAGATTGAGTGAAGCTGAGATTCGACAGTGAAGCCGAATCAC 4545
QY 288 AGATCAGCAGAGTGAAGAGT 308
DB 4544 GGGCAACGAGTGAATCTGT 4524

RESULT 22
AAS28363/c
ID AAS28363 standard; DNA; 32146 BP.
XX
XX AAS28363;
AC
XX
DT 07-NOV-2001 (first entry)
XX
DE Genomic sequence #203 encoding for novel human respiratory antigen.
XX
KW Human; respiratory antigen; respiratory disorder; throat disorder;
lung disorder; nose disorder; lung cancer; gene therapy; cytostatic;

KW anti allergic; anti asthmatic; anti inflammatory; olfactory;
XX respiratory active; ds.
XX Homo sapiens.
XX WO20015448-A1.
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XX 02-AUG-2001.
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XX 17-JAN-2001; 2001WO-US001333.
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XX 31-JAN-2000; 2000US-0179065P.
PR 04-FEB-2000; 2000US-0180628P.
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 PR 27-SEP-2000; 2000US-0235200P.

PR 05-JAN-2001; 2001US-0259678P.
 XX (HUMA-) HUMAN GENOME SCI INC.
 PA Rosen CA, Barash SC, Ruben SM;
 PI WPI; 2001-476224/51.
 XX
 DR
 XX
 PT Isolated polypeptide for treating, preventing and/or prognosing
 PT disorders related to the respiratory system including respiratory cancers
 PT and also for testing and detection e.g. diagnosis.
 XX
 PS
 XX Disclosure; SEQ ID NO 797; 546bp; English.
 CC The present invention relates to the isolation of novel human respiratory
 CC antigens (AAU17685-AAU17975), and cDNA and genomic sequences encoding for
 CC these polypeptides. The sequences of the invention are useful for
 CC preventing, treating and/or prognosing disorders related to the
 CC respiratory system including throat disorders (e.g. vocal cord paralysis,
 CC tonsillitis, and laryngitis), lung disorders e.g. pneumonia, allergic
 CC disorders e.g. asthma, pleurisy, cystic fibrosis, emphysema, nose
 CC disorders and cancers of the respiratory tissues e.g. lung cancer. The
 CC polynucleotide sequences of the invention are useful in gene therapy and
 CC antisense therapy. AAU17685-AAU17975 represent genomic sequences encoding
 CC for novel human respiratory antigens. Note: The sequence data for this
 CC patent did not form part of the printed specification, but was obtained
 CC in electronic format directly from WIPO at
 CC ftp.wipo.int/pub/published_pct_sequences
 XX
 SQ Sequence 32146 BP; 8745 A; 6747 C; 7148 G; 9506 T; 0 U; 0 Other;
 Query Match 37.7%; Score 150.6; DB 4; Length 32146;
 Best Local Similarity 68.5%; Pred. No. 1.9e-31;
 Matches 224; Conservative 1; Mismatches 95; Indels 7; Gaps 1;
 QY 81 ATGCTGTAATCCGACGACCTTCGGAGGCAAGTGGGGGATCACTGAGGTCAAGAGA 140
 DB 2437 ACGCTGTAACTCTGACCTTTGGAGGCGGAGGATCACTTGAAGTCAAGAGT 2378
 QY 141 TCGAGACCACTCTGGCCAAATGTAAGCCCGTCTTACTTAAATCAAAAAATAGC 200
 DB 2377 TTGAGACCAAGCTGCGCAATGTAAGCCCGTCTTACTTAAATCAAAAAATAGC 2318
 QY 201 TGGGATGTTGGACACACCTGTATCCCACTACTCAGAG-----CCGAGATTGC 253
 DB 2317 AGGGCATGTGCGGGACCTGTATCCCACTGTGGAGACTGAGCAGAGAGATTG 2258
 QY 254 AGTGAGCTGATGTCGACAGTGAAGCAATCACTACAGAGTGAAGTGAAGAGC 313
 DB 2257 CTTGAACCCGGAGGCGAGGTTGCAATGAGCCGAGTTGACTGCAATGCAAGAGC 2198
 QY 314 KCCGTCTCAAAAACAACAAAAAACAACCAATAGCATTTGTCATCTCGGTT 373
 DB 2197 TCCGTTCAAAAAAAGATTAAGTGAAGCAATAGAGGGA 2138
 QY 374 CCCAGACTATTGACAGAGACCAAAAG 400
 DB 2137 TCTCAGCAAAATCTGATTAAACAAAG 2111
 RESULT 23
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 ID ADG41559 standard; DNA; 32146 BP.
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 XX ADG41559;
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 DT 26-FEB-2004 (first entry)
 XX
 XX Human respiratory system associated genomic DNA seq id 797.
 DE
 XX Human respiratory system associated genomic DNA seq id 797.
 DE
 XX antiinflammatory; antiallergic; antiasmatic; cyostatic; gene therapy;
 KW respiratory system antigen;
 KW human respiratory system associated polynucleotide;

PR 08-DEC-2000; 2000US-0251868P.
 PR 08-DEC-2000; 2000US-0251869P.
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 PR 08-DEC-2000; 2000US-0251990P.
 PR 11-DEC-2000; 2000US-0254097P.
 PR 05-JAN-2001; 2001US-0259678P.
 PR 17-JAN-2001; 2001US-00764860.
 PR 14-FEB-2002; 2002US-00074095.

XX (HUMA-) HUMAN GENOME SCI INC.
 XX
 XX Rosen CA, Ruben SM, Barash SC,
 XX
 XX WPI; 2003-902033/82.

XX Novel respiratory system antigen and polynucleotides encoding the
 PT polypeptides; useful for treating, diagnosing, treating or preventing
 PT conellitis, pneumonia, asthma and cystic fibrosis, emphysema, throat
 PT cancer.

XX Discloure; SEQ ID NO 797; 236pp; English.

XX The invention describes an isolated polypeptide (I) comprising an amino
 CC acid sequence that is at least 90% identical to polypeptide fragment of
 CC any one of 299 respiratory system antigen sequences (PS) and having
 CC biological activity, polypeptide domain or epitope of PS, full-length
 CC protein of PS or variant, allelic variant or species homolog of PS. (I)
 CC or a polynucleotide (II) encoding (I) is also useful for diagnosing a
 CC pathological condition or a susceptibility to a pathological condition in
 CC a subject which involves determining the presence or absence of mutation
 CC in (II) or determining the presence or amount of expression of (I) in a
 CC biological sample and diagnosing a pathological condition based on the
 CC result. The human respiratory system associated polynucleotides, the
 CC polypeptides encoded by them, and antibodies that immunospecifically bind
 CC these polypeptides are useful in diagnosis, treatment, prevention and/or
 CC prognosis of disorders of respiratory system such as throat disorders
 CC (e.g., vocal cord paralysis, tonsillitis, and laryngitis), lung disorders
 CC (e.g., pneumonia), allergic disorders (e.g., asthma and eosinophilic
 CC pneumonia), pleurisy, cystic fibrosis, emphysema, histiocytosis,
 CC sarcoidosis, nose disorders (rhinitis and sinusitis), neoplasms and/or
 CC cancers of respiratory tissues (e.g., throat cancer, lung cancer, and
 CC cancer of the nose). The polynucleotides are useful in gene therapy
 CC techniques, for chromosome identification, identifying individuals from

Query Match 37.7%; Score 150.6; DB 10; Length 32146;

Best Local Similarity 68.5%; Pred. No. 1.9e-31;
 Matches 224; Conservative 1; Mismatches 95; Indels 7; Gaps 1;

XX 81 ATGCTGTATCCGACGACCTTGGAGGCGAAGTGGGCGATCACTGAGGTCAAGGA 140
 DB 2437 ACGCTGTATGCTATGACCTTTGGAGGCGAAGTGGGCGATCACTGAGGTCAAGGA 2378
 XX 141 TCGAGACCATCTGCGCAACATGTGAACCCGCTCTTACTAATAAATCAAAAATATAGC 200
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 XX 201 TGGGATGTGGGACACACTGTAGTCCGAGTACTACAGAG-----CCGAGATTGC 253
 DB 2317 AGGGATGTGGGCGGACCTGTAGTCCGAGTACTGTGGGAGACTGAGCCAGAGAAATTG 2258
 XX 254 AGTGAGTGAGATGCGAGAGTGAAGCGAAATCAAGATCAAGAGTGAAGAGAGAC 313
 DB 2257 CTTGAACCCGGAGGCGGAGGTGCAATGAGCCGAGATTGTACTGCAATGCAAGAGAC 2198
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 DB 2137 TCTCAGCAAAATCTGATTTAACAAG 2111

RESULT 24
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 XX 04-NOV-2004 (first entry)
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 XX Human respiratory system associated polypeptide-related DNA SeqID797.
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 XX respiratory system-related polypeptide; antiasthmatic; antibacterial;
 KW antiinflammatory; cytoskeletal; antianaemic; antiallergic; gene therapy;
 KW pneumonia; lung cancer; cystic fibrosis; asthma; sarcoidosis; rhinitis;
 KW anaemia; leukaemia; inflammation; sinusitis;
 KW chronic obstructive pulmonary disease; infectious disease; human; ds.
 OS
 XX Homo sapiens.
 XX
 XX US2003077704-A1.
 PN
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 XX 24-APR-2003.
 PD
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 XX 14-FEB-2002; 2002US-00074095.
 PF
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 XX 31-JAN-2000; 2000US-0179065P.
 PR 04-FEB-2000; 2000US-0180628P.
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 PR 30-AUG-2000; 2000US-0228924P.
 PR 01-SEP-2000; 2000US-0229287P.
 PR 01-SEP-2000; 2000US-0229343P.
 PR 01-SEP-2000; 2000US-0229344P.
 PR 05-SEP-2000; 2000US-0229345P.
 PR 05-SEP-2000; 2000US-0229509P.
 PR 06-SEP-2000; 2000US-0229513P.
 PR 06-SEP-2000; 2000US-0230437P.
 PR 08-SEP-2000; 2000US-0230438P.
 PR 08-SEP-2000; 2000US-0231242P.
 PR 08-SEP-2000; 2000US-0231243P.
 PR 08-SEP-2000; 2000US-0231244P.
 PR 08-SEP-2000; 2000US-0231413P.
 PR 08-SEP-2000; 2000US-0231414P.

ID	ADL13941/c	ADL13941 standard; DNA, 125515 BP.
XX	ADL13941;	
XX	06-MAY-2004 (first entry)	
XX	Osteoarthritis-associated polymorphic nucleotide #473.	
XX	ds; gene; osteopathic; antiinflammatory; antiarthritic; gene therapy;	
XX	joint space narrowing; osteophyte development; joint pain;	
XX	osteoarthritis; SNP, single nucleotide polymorphism.	
XX	Homo sapiens.	
XX	MO2003054166-A2.	
XX	03-JUL-2003.	
XX	19-DEC-2002; 2002MO-US041225.	
XX	20-DEC-2001; 2001US-0342603P.	
XX	(INCY-) INCYTE GENOMICS INC.	
XX	Jones KA, Schafer A;	
XX	WPI; 2003-559141/52.	
XX	Determining susceptibility of an individual to joint space narrowing,	
XX	osteoarthritis development and/or joint pain comprises identifying whether	
XX	the individual has at least one polymorphism in a polymorphic nucleotide encoding	
XX	a protein.	
XX	Disclosure; SEQ ID NO 473; 297bp; English.	
XX	The invention relates to a method of determining susceptibility of an	
XX	individual to joint space narrowing and/or osteophyte development and/or	
XX	joint pain comprising identifying whether the individual has at least one	
XX	polymorphism in a polymorphic nucleotide encoding at least one of the protein	
XX	listed in the specification. The methods, composition and agent are	
XX	useful for modulating the susceptibility of an individual to joint space	
XX	narrowing and/or osteophyte development and/or joint pain that is	
XX	associated with a disease, preferably osteoarthritis. The cell line and	
XX	the non-human animal are useful for screening for an agent for diagnosing	
XX	an individual having susceptibility to joint space narrowing and/or	
XX	osteoarthritis development and/or joint pain. This sequence corresponds to	
XX	the polymorphic nucleotide encoding a protein listed in the specification. (Note:	
XX	The sequence data for this patent did not form part of the printed	
XX	specification but was obtained in electronic format directly from WIPO at	
XX	ftp.wipo.int/pub/published_pct_sequences).	
XX	Sequence 125515 BP; 33180 A; 28822 C; 28744 G; 34769 T; 0 U; 0 Other;	
XX	Query March 37.7%; Score 150.6; DB 10; Length 125515;	
XX	Best Local Similarity 75.7%; Pred. No. 3e-31;	
XX	Matches 202; Conservative 1; Mismatches 55; Indels 9; Gaps 1;	
XX	81 ATGCGCTTAATCCAGCACTTCGGGAGGCGCAAGGTGGGCGGATCACCTGAGGTCAGAGAGA 140	
XX	Db ATGCTTTAATCCAGCACTTCGGGAGGCGTGAAGAGGCGGATTAATTGAGGTCAAGACT 5274242	
XX	141 TCGAGACCAATCTGGCCACAATGGTGAACCCCGTCTTTACTTAAATAATCAAAAATTAC 200	
XX	Db TTCAGACCAAGCTGGCCACAATGGTGAACCCCGTCTTAAATAATCAAAAATTAC 526822	
XX	201 TGGGCAATGGGCGCACACCTGTAGTCCGAGCTACACAGAGCGGAGATTGAGAGAAC 260	
XX	Db CGGGCTGGTGGAGCACCACTGTAGTCCGAGCTACACAGAGCACAGAGTTGAGAGAAC 526222	

Qy	261	TGAGTGCAGAGTGAGCCGAATTCACAGTTCACAGAGTGAGTGACKCGCT	320
Db	52621	CAGATATATCCCACTGCATCTCCAGCTCGGCGCA-----CAGAGCGAGCTCCACT	52577
Qy	321	CAAAAACACACACAAAAAACAACAAAAA	347
Db	52570	CAAAAAAAAAAAAAAAAAAAAAAG	52544
RESULT 26			
ID	AAZ29204/C		
XX	AAZ29204 standard; DNA; 17538 BP.		
XX	AAZ29204;		
XX	21-FEB-2000 (first entry)		
XX	Human myelin oligodendrocyte glycoprotein gene.		
XX	Human: myelin oligodendrocyte glycoprotein; MOG: NS-specific antigen;		
KW	nervous system-specific antigen; T cell; peripheral nervous system; PNS;		
KW	central nervous system; CNS; nerve regeneration; neuronal degeneration;		
KW	spinal cord injury; blunt trauma; penetrating trauma; senile dementia;		
KW	ischemic stroke; diabetic neuropathy; glaucoma; haemorrhagic stroke;		
KW	Alzheimer's disease; Parkinson's disease; Huntington's chorea;		
KW	amyotrophic lateral sclerosis; ALS; treatment; ds.		
XX	Homo sapiens.		
OS			
XX	Key	Location/Qualifiers	
XX	FT CDS	1166..15142	
FT		/trac= a	
FT		/product= "myelin oligodendrocyte glycoprotein"	
FT		/note= "reading frame is interrupted by introns the	
FT		precise location of which is not given in the	
FT		specification "	
XX			
XX	MO9960021-A2.		
XX	25-NOV-1999.		
XX	19-MAY-1999;	99MO-US010953.	
XX	19-MAY-1998;	98IL-00124550.	
PR	21-JUL-1998;	98MO-US014715.	
PR	22-DEC-1998;	98US-00218277.	
XX	(YEDA) YEDA RES & DEV CO LTD.		
XX	(MCIN/) MCINNIS P A.		
XX	Eisenbach-Schwartz M, Cohen IR, Beserman P, Mosonogo A, Moalem G;		
XX	WPI; 2000-072430/06.		
XX	P-PSDB; AAY44236.		
XX	New compositions useful to treat nervous system injury or disease e.g.		
XX	traumatic injury, Alzheimer's disease etc.		
XX	Claim 9; Fig 18; 92pp; English.		
XX	The present sequence is a gene encoding human myelin oligodendrocyte		
XX	glycoprotein which is a nervous system-specific antigen. The antigen or		
XX	peptides derived from it activate T cells in vivo. The present sequence		
XX	is used to promote nerve regeneration or to prevent or inhibit neuronal		
XX	degeneration caused by injury or diseases of nerves within the CNS or		
XX	PNS. Such injury includes spinal cord injury, blunt trauma, penetrating		
XX	trauma, haemorrhagic stroke or ischaemic stroke, whilst diseases include		
XX	diabetic neuropathy, senile dementia, Alzheimer's disease, Parkinson's		
XX	disease, glaucoma, Huntington's chorea, amyotrophic lateral sclerosis,		
XX	etc		
Qq	Sequence 17538 BP; 4624 A; 4121 C; 3990 G; 4803 T; 0 U; 0 Other;		

Query Match 37.6%; Score 150.4; DB 3; Length 17538;
 Best Local Similarity 72.1%; Pred. No. 1.8e-31;
 Matches 196; Conservative 0; Mismatches 76; Indels 0; Gaps 0;

QY 81 ATGCTGTAAATCCAGACCTTCGGAGGCGCAAGGTGGGCGGATCACTGAGGTCAAGAGA 140
 DB 13528 ATGCTGTAAATCCAGACCTTCGGAGGCGCAAGGTGGGCGGATCACTGAGGTCAAGAGA 13469
 QY 141 TCGAGACATCTCTGGCCCAACATGTGTAACCCCGCTTACTTAAATAACAAAAATATGC 200
 DB 13468 TCAAGACCCAGGCTTGCCCAACATGTGTAACCCCGCTTACTTAAATAACAAAAATATGC 13409
 QY 201 TGGGCATGTGGACACACCTGTAGTCCCACTACTCAAGACCGGAGATTGACGTAGC 260
 DB 13408 TGGGCATGTGGACATGGCCCTGTAGTTCACGACCTTGGAGGCTTGAGGCAAGAAATCG 13349
 QY 261 TGAAGTCGACAGTGAAGCCGAATATCAAGATCAAGATGAGCAGAGTGAAGACKCCGCT 320
 DB 13348 CTTGAACCCAGAGGTGAGGTTGCAGTGAGCCGAGATTGTGCCATTCATCTCA 13289
 QY 321 CAAAAACAACAACAAAAACAAAAACCATTA 352
 DB 13288 AAAAAACTTCACTCCAAAAAAGAAAGAAA 13257

RESULT 27
 ABN86597/C
 ID ABN86597 standard; DNA; 17538 BP.
 AC ABN86597;
 XX
 XX
 DT 05-NOV-2002 (first entry)

Human myelin oligodendrocyte glycoprotein (MOG) gene sequence.

XX
 XX
 DB Nerve regeneration; neuroprotection; neuronal degeneration; CNS; PNS;
 XX central nervous system; peripheral nervous system; tranquillizer; MOG;
 XX vulnerability; cerebroprotective; anti-tumour; antidiabetic; anticonvulsant;
 XX neurotrophic; antiparkinsonian; ophthalmological; analgesic; hepatotropic;
 XX osteopathic; vasotropic; nephrotropic; cytostatic; antigen; gene therapy;
 XX myelin oligodendrocyte glycoprotein; human; gene; ds.
 OS
 XX Homo sapiens.
 XX
 XX US2002072493-A1.
 XX
 PD 13-JUN-2002.
 XX
 XX 28-JUN-2001; 2001US-00893348.
 PF
 XX 19-MAY-1998; 98IL-00124500.
 PR 21-JUL-1998; 98MO-US014715.
 PR 22-DEC-1998; 98US-00218277.
 PR 19-MAY-1999; 99US-00314161.
 XX
 XX (YEDA) YEDA RES & DEV CO LTD.
 XX
 XX Eisenbach-Schwartz M, Hauben E, Cohen IR, Beeserman P, Mosonogo A;
 PI Moalem G;
 XX
 XX WPI; 2002-607255/65.
 DR P-PSDB; ABB81071.
 XX
 XX Promoting nerve regeneration and preventing neuronal degeneration in the
 PT central/peripheral nervous system from injury/disease, comprises
 PT administering nervous system-specific activated T cells/antigen, or
 PT analogs/peptides.
 XX
 XX
 PS Disclosure; Fig 18; 93pp; English.
 CC The invention relates to promoting nerve regeneration or conferring
 CC neuroprotection and preventing or inhibiting neuronal degeneration in the
 CC central/peripheral nervous system (NS). The method involves administering

CC NS-specific activated T cells, NS-specific antigen, its analogue or its
 CC peptide, a nucleotide sequence the NS-specific antigen or its analogue or
 CC combinations. The method is useful for promoting nerve regeneration and
 CC preventing neuronal degeneration in central/peripheral nervous system
 CC from injury/disease, where the injury is spinal cord injury, blunt
 CC trauma, penetrating trauma, hemorrhagic stroke, ischemic stroke or
 CC damages caused by surgery such as tumour excision. The disease is not an
 CC autoimmune disease or neoplasia. The disease results in a degenerative
 CC process occurring in either gray or white matter or both. The disease is
 CC diabetic neuropathy, senile dementia, Alzheimer's disease, Parkinson's
 CC disease, facial nerve (Bell's) palsy, glaucoma, Huntington's chorea,
 CC amyotrophic lateral sclerosis, non-arteritic optic neuropathy, and
 CC vitamin deficiency, intervertebral disc herniation, prion diseases such
 CC as Creutzfeldt-Jakob disease, carpal tunnel syndrome, peripheral
 CC neuropathies associated with various diseases, including but not limited
 CC to uremia, porphyria, hypoglycemia, Sjogren-Larson syndrome, acute
 CC sensory neuropathy, chronic ataxic neuropathy, biliary cirrhosis, primary
 CC amyloidosis, obstructive lung diseases, acromegaly, malabsorption
 CC syndromes, polycythemia vera, immunoglobulin (Ig)A- and IgG gamma-
 CC pathies, complications of various drugs (e.g., metronidazole) and toxins
 CC (e.g., alcohol or organophosphates), Charcot-Marie-Tooth disease, ataxia
 CC telangiectasia, Friedreich's ataxia, amyloid polynuropathies,
 CC adrenomyeloneuropathy, Giant axonal neuropathy, Refsum's disease, Fabry's
 CC disease, or lipoproteinemia. The present sequence represents a human
 CC myelin oligodendrocyte glycoprotein (MOG) gene sequence, an example of NS
 CC -specific antigen

XX
 XX
 SQ Sequence 17538 BP; 4624 A; 4120 C; 3991 G; 4803 T; 0 U; 0 Other;

Query Match 37.6%; Score 150.4; DB 6; Length 17538;
 Best Local Similarity 72.1%; Pred. No. 1.8e-31;
 Matches 196; Conservative 0; Mismatches 76; Indels 0; Gaps 0;

QY 81 ATGCTGTAAATCCAGACCTTCGGAGGCGCAAGGTGGGCGGATCACTGAGGTCAAGAGA 140
 DB 13528 ATGCTGTAAATCCAGACCTTCGGAGGCGCAAGGTGGGCGGATCACTGAGGTCAAGAGA 13469
 QY 141 TCGAGACATCTCTGGCCCAACATGTGTAACCCCGCTTACTTAAATAACAAAAATATGC 200
 DB 13468 TCAAGACCCAGGCTTGCCCAACATGTGTAACCCCGCTTACTTAAATAACAAAAATATGC 13409
 QY 201 TGGGCATGTGGACACACCTGTAGTCCCACTACTCAAGACCGGAGATTGACGTAGC 260
 DB 13408 TGGGCATGTGGACATGGCCCTGTAGTTCACGACCTTGGAGGCTTGAGGCAAGAAATCG 13349
 QY 261 TGAAGTCGACAGTGAAGCCGAATATCAAGATCAAGATGAGCAGAGTGAAGACKCCGCT 320
 DB 13348 CTTGAACCCAGAGGTGAGGTTGCAGTGAGCCGAGATTGTGCCATTCATCTCA 13289
 QY 321 CAAAAACAACAACAAAAACAAAAACCATTA 352
 DB 13288 AAAAAACTTCACTCCAAAAAAGAAAGAAA 13257

RESULT 28
 ADQ97695/C
 ID ADQ97695 standard; DNA; 88892 BP.
 AC ADQ97695;
 XX
 XX
 DT 07-OCT-2004 (first entry)
 XX
 XX Human cancer associated sequence HD10-029, SEQ ID 672.
 DE
 XX
 XX Cytostatic; Gene Therapy; cancer; leukemia; lymphoma; Human; ds.
 KW
 XX
 XX Homo sapiens.
 OS
 XX
 XX WO2004060304-A2.
 XX
 XX
 XX 22-JUL-2004.
 PD
 XX 22-DEC-2003; 2003WO-US041389.
 PF

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XX 27-DEC-2002; 2002US-00330773.
XX (SAGR-) SAGRES DISCOVERY INC.
XX Morris DW, Malandro MS;
XX WPI; 2004-543781/52.
XX
XX New isolated cancer associated nucleic acids comprising at least 10
XX contiguous nucleotides, useful for diagnosing, preventing and/or treating
XX cancers such as leukemia and lymphoma.
XX
XX Claim 1; SEQ ID NO 672; 1999p; English.
XX
XX The present invention relates to cancer associated sequences (ADQ97025-
XX ADQ98004). The sequences are useful for the diagnosis, prevention and/or
XX treatment of cancer, such as leukemia and lymphoma. Note: The sequence
XX data for this patent did not form part of the printed specification, but
XX was obtained in electronic format directly from WIPO at
XX ftp.wipo.int/pub/published_pct_sequences.
XX
XX Sequence 88892 BP; 25748 A; 16788 C; 17057 G; 27074 T; 0 U; 2225 Other;
XX
XX Query Match 37.6%; Score 150.4; DB 12; Length 88892;
XX Best Local Similarity 74.5%; Pred. No. 3e-31;
XX Matches 204; Conservative 1; Mismatches 62; Indels 7; Gaps 1;
XX
XX 81 ATGCTGTAAATCCAGCACTTCGGAGAGCCAAAGTGGCGGATCACTGAGGTCAAGAGA 140
XX 58516 ACGCTGTAAATCCAGCACTTCGGAGAGCCAAAGTGGCGGATCACTGAGGTCAAGAGA 140
XX 141 TCGAGACCATCTGGCCCAATGATGTGAACCCCGTCTTAAATAAATACAAAATAGC 200
XX 58456 TCGAGACCATCTGGCCCAATGATGTGAACCCCGTCTTAAATAAATACAAAATAGC 200
XX 201 TGGGATGATGGGACACAGCTGTACTCCAGTACTTCAAGG-----GCCGGAGATTGC 253
XX 58396 TGGGATGATGGGACACAGCTGTACTCCAGTACTTCAAGGAGCTGAGGAGGAAATTG 58337
XX 254 AGTAGCTGAGATCGCAGAGTGAAGCCGAATATCAGATCAAGAGTGAAGTGAAGC 313
XX 58336 CTGTAATAGAGCTGAGATCAATGCCACTGCACTCCAGGCTCAGCGCAAGAGTGAAGC 58277
XX 314 KCCGCTTCAAAAACAACAACAAAACAAAACAAA 347
XX 58276 TTTGTCTCAAAAAAAGAAAAAAGAAAAA 58243
XX
XX RESULT 29
XX AAI61042 standard; cDNA; 4779 BP.
XX AAI61042;
XX
XX 22-OCT-2001 (first entry)
XX
XX Human polynucleotide SEQ ID NO 5031.
XX
XX Human; nocturnal; immunosuppressant; cytosolic; gene therapy; cancer;
XX peripheral nervous system; neuropathy; central nervous system; CNS;
XX Alzheimer's; Parkinson's disease; Huntington's disease; haemostatic;
XX amyotrophic lateral sclerosis; Shy-Drager Syndrome; chemotactic;
XX chemokine; thrombolytic; drug screening; arthritis; inflammation;
XX leukaemia; ss.
XX
XX Homo sapiens.
XX
XX WO200153312-A1.
XX
XX 26-JUL-2001.
XX
XX 26-DEC-2000; 2000WO-US034263.

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XX 23-DEC-1999; 99US-00471275.
XX 21-JAN-2000; 2000US-00488725.
XX 25-APR-2000; 2000US-00552317.
XX 20-JUN-2000; 2000US-00598042.
XX 19-JUL-2000; 2000US-00620312.
XX 03-AUG-2000; 2000US-00653450.
XX 14-SEP-2000; 2000US-00662191.
XX 19-OCT-2000; 2000US-00693036.
XX 29-NOV-2000; 2000US-00727344.
XX
XX (HYSE-) HYSEQ INC.
XX
XX Tang YT, Liu C, Asundi V, Chen R, Ma Y, Qian XB, Ren F, Wang D,
XX Wang J, Wang Z, Wehrman T, Xu C, Xue AJ, Yang Y, Zhang J, Zhao QA,
XX Zhou P, Goodrich R, Dymanac RT;
XX
XX WPI; 2001-442253/47.
XX P-P8DB; AAM41886.
XX
XX Novel nucleic acids and polypeptides, useful for treating disorders such
XX as central nervous system injuries.
XX
XX Claim 1; SEQ ID NO 5031; 10078bp; English.
XX
XX The invention relates to human nucleic acids (AA157798-AA161369) and the
XX encoded polypeptides (AAM38642-AAM42213) with nocturnal,
XX immunosuppressant and cytosolic activity. The polynucleotides are useful
XX in gene therapy. A composition containing a polypeptide or polynucleotide
XX of the invention may be used to treat diseases of the peripheral nervous
XX system, such as peripheral nervous injuries, peripheral neuropathy and
XX localized neuropathies and central nervous system diseases, such as
XX Alzheimer's, Parkinson's disease, Huntington's disease, amyotrophic
XX lateral sclerosis, and Shy-Drager Syndrome. Other uses include the
XX utilization of the activities such as: Immune system suppression,
XX Activin/inhibin activity, chemotactic/chemokinetic activity, hemostatic
XX and thrombolytic activity, cancer diagnosis and therapy, drug screening,
XX assays for receptor activity, arthritis and inflammation, leukemias and
XX C.N.S disorders. Note: The sequence data for this patent did not form
XX part of the printed specification
XX
XX Sequence 4779 BP; 1453 A; 960 C; 1048 G; 1318 T; 0 U; 0 Other;
XX
XX Query Match 37.5%; Score 150; DB 4; Length 4779;
XX Best Local Similarity 79.7%; Pred. No. 1.6e-31;
XX Matches 177; Conservative 0; Mismatches 45; Indels 0; Gaps 0;
XX
XX 81 ATGCTGTAAATCCAGCACTTCGGAGAGCCAAAGTGGCGGATCACTGAGGTCAAGAGA 140
XX 4253 ACGCTGTAAATCCAGCACTTCGGAGAGCCAAAGTGGCGGATCACTGAGGTCAAGAGA 140
XX 141 TCGAGACCATCTGGCCCAATGATGTGAACCCCGTCTTAAATAAATACAAAATAGC 200
XX 4313 TCAAGACCATCTGGCCCAATGATGTGAACCCCGTCTTAAATAAATACAAAATAGC 200
XX 201 TGGGATGATGGGACACAGCTGTACTCCAGTACTTCAAGGAGCTGAGGAGGAAATTG 260
XX 4373 TGGGATGATGGGACACAGCTGTACTCCAGTACTTCAAGGAGCTGAGGAGGAAATTG 260
XX 261 TGAATGCGCAGAGTGAAGCCGAATATCAGATCAAGAGTGAAG 302
XX 4433 CGAGATTGGCGCACTGCACTCCAGCTGGGCGGAGCAGAGTAG 4474
XX
XX RESULT 30
XX AED18472 standard; DNA; 100998 BP.
XX AED18472;
XX
XX 15-DEC-2005 (first entry)
XX
XX Fibrotic disorder associated polynucleotide SEQ ID NO 723.

```

XX antiinflammatory; gene therapy; fibrogenesis; gene expression;
KM therapeutic; diagnosis; uterine fibroids; gynecological; inflammation;
KM ds.
XX Homo sapiens.
XX MO2005098041-A2.
XX
XX 20-OCT-2005.
XX
XX 28-MAR-2005; 2005WO-US010257.
XX
XX 26-MAR-2004; 2004US-0556546P.
XX 19-OCT-2004; 2004US-0620444P.
XX 15-DEC-2004; 2004US-0636240P.
XX
XX (UYFL) UNIV FLORIDA RES FOUND INC.
XX
XX Chegini N, Luo X, Ding L, Williams RS.
XX
XX WPI; 2005-703565/72.
XX
XX Identifying a modulator of a gene that is differentially-expressed in
XX fibrotic tissue or during fibrogenesis, or a polypeptide encoded by the
XX gene, in a cell population by contacting the cell population with a test
XX agent.
XX
XX Disclosure; SEQ ID NO 723; 202pp; English.
XX
XX The invention describes a method of identifying a modulator of at least
XX one gene that is differentially-expressed in fibrotic tissue or during
XX fibrogenesis, or a polypeptide encoded by the differentially-expressed
XX gene, in a cell population, comprising contacting the cell population
XX with a test agent, and determining if the test agent modulates the
XX expression of the gene or biological activity of the polypeptide encoded
XX by the gene. Also described are: detecting a fibrotic disorder in a
XX subject; modulating gene expression in fibrotic tissue; and an array
XX comprising a substrate having addresses, where each address has a capture
XX probe that can specifically bind at least one polynucleotide that is
XX differentially expressed in fibrotic disorders, or its complement. The
XX method is useful in identifying a modulator of at least one gene that is
XX differentially-expressed in fibrotic tissue or during fibrogenesis, or a
XX polypeptide encoded by the differentially-expressed gene, in a cell
XX population for preparing a composition for diagnosing or treating
XX fibrotic disorders, e.g. uterine fibrosis. This sequence represents a
XX polynucleotide associated with detection and treatment of fibrotic
XX disorders. Note: This sequence does not appear in the printed
XX specification but has been obtained in electronic format directly from
XX WIPO at ffp.wipo.int/pub/published_pct_sequences.
XX
XX Sequence 100998 BP; 25449 A; 22644 C; 23083 G; 29822 T; 0 U; 0 Other;
XX
XX Query Match 37.5%; Score 150; DB 14; Length 100998;
XX Best Local Similarity 69.4%; Pred. No. 4,1e-31;
XX Matches 225; Conservative 1; Mismatches 81; Indels 17; Gaps 1;
XX
XX 30 GGAACCCAAATATTAATAGACATTGTCAGGCCAGGCATGACATGGCTGAATGCCCTGTA 89
XX 92984 GAAACCCCTCTCTACTAAAGTACAAAAAATTAAGCCGGCGCTGGGCGGCTGTA 93043
XX
XX 90 ATCCCAAGCCTTCGGGAGGCCAAGTGGCGCATCATCCTGAGTCAAGAGATCGAGACCA 149
XX 93044 ATCCCAAGCCTTCGGGAGGCCAAGGAGGTGATCACTGAGGTCAAGGTTCAAGACCA 93103
XX
XX 150 TCCTGGCCAACTGGGAACCCCGTCTTACTTAATAATCAAAAAATAGCTGGCGATGG 209
XX 93104 GCTTGCCCAACTGGGAACCCCGTCTTACTTAATAATCAAAAAATAGCTGGCGATGG 93163
XX
XX 210 TGGCAACACCTGTAGTCCAGCTACTCAGAGCCGAGATTGCACTGAGTGAATGCC 269
XX 93164 TGGCAACACCTGTAGTCCAGCTACTCAGAGCCGAGATTGCACTGAGTGAATGCC 93223

QY 270 AGAGTGAGCCGAATATCACAGATCA-----CAGAGTGACAGAGTGAGA 312
DB 93224 AGAGCTGCAAGTATGAGCCGAGATATCAGCCACTGACCTAGCTGGGTGATGAGAGTGAAA 93283
QY 313 CKCGTCTCAAAAACAAACAACAA 336
DB 93284 CTCATCTCAACAACAACAACAA 93307
RESULT 31
AEB96535/c
ID AEB96535 standard; DNA; 151909 BP.
XX
XX AEB96535;
XX
XX AC
XX 06-OCT-2005 (first entry)
XX
XX Human CABIN1 gene, SEQ ID 19.
XX
XX hepatitis C virus infection; antiinflammatory; hepatotropic; virucide;
KM liver cirrhosis; fibrosis; hepatoma; SNP detection; CABIN1; ds.
XX
XX Homo sapiens.
XX
XX Location/Qualifiers
XX 7588
XX Key
XX variation
XX
XX location
XX /tag= a
XX /standard_name= "Single nucleotide polymorphism"
XX
XX location
XX /tag= b
XX /standard_name= "Single nucleotide polymorphism"
XX
XX location
XX /tag= c
XX /standard_name= "Single nucleotide polymorphism"
XX
XX location
XX /tag= d
XX /standard_name= "Single nucleotide polymorphism"
XX
XX location
XX /tag= e
XX /standard_name= "Single nucleotide polymorphism"
XX
XX location
XX /tag= f
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XX location
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XX location
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XX location
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XX location
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XX /standard_name= "Single nucleotide polymorphism"
XX
XX location
XX /tag= p
XX /standard_name= "Single nucleotide polymorphism"
XX

QY 144 AGACCATCTTGGCCCAACATGTTGAACCCCGTCTTTACTATAAATAACAAAATATAGCTGG 203
DB 527 AGACCCAGCTTGGCCCAACATGTTGAACCCCGTCTTTACTATAAATAACAAAATATAGCTGG 468
QY 204 GCATGTGTGGCAGACACCTGTAGTCCAGCTACTCAGAGCCGAGATTCAG--TGAGCT 261
DB 467 GCATGTGTGGCAGATGTGCCCTATATCTCCAGCTACTTGGGAAGCTGAGGCGAGGAATCGCTT 408
QY 262 GAGATTCGAGAGTGAAGCCGAATATCAGATCAAGAGTGAAGAGTGAACCKCCGCTTC 321
DB 407 GAACCTGGGAGGCGAGAGCCACTGCACCTCGGGGCAACAGAGTGAAGGCTGTCTC 348
QY 322 AAAAACAACAACAAAAAACAATAAGCATGTGCTCATCTGCGTCCAG 378
DB 347 CAAAAACAAAAACAAAAACAAAAACAAAGATTAACCCAAAGGAGATGCAAG 291

RESULT 32

ACN44170
ID ACN44170 standard; DNA, 196686 BP.

ACN44170;

18-NOV-2004 (first entry)

Human genomic sequence hCG39530.

Cytostatic; carcinoma; lymphoma; cancer; human; gene; ss.

Homo sapiens.

WO2003073826-A2.

12-SEP-2003.

28-FEB-2003; 2003WO-US006235.

01-MAR-2002; 2002US-00087192.

(SAGR-) SAGRES DISCOVERY.

Morris DW;

WPI; 2003-328604/31.

Recombinant nucleic acid useful for diagnosis and treatment of carcinoma comprises a nucleotide sequence.

Claim 1; SEQ ID NO 484; Opp: English.

The present invention relates to novel DNA and protein sequences which are associated with carcinomas. The sequences are useful for: (i) for screening drug candidates; (ii) for screening of bioactive agent capable of binding to Carcinoma Associated Protein (CAP); (iii) for screening of a bioactive agent capable of modulating the activity of CAP; (iv) for evaluating the effect of a candidate carcinoma drug; (v) for diagnosing carcinoma; (vi) for inhibiting the activity of CAP; (vii) for treating carcinoma; (viii) for neutralizing the effect of CAP; (ix) as a biochip; (x) for diagnosing carcinoma or a propensity to carcinoma; and (xi) for determining Carcinoma Associated (CA) gene copy number. In addition, the CA genes are useful as DNA vaccines and the CAP are useful as markers of carcinoma including lymphoma. The present sequence is one such CA coding sequence. Note: This patent is an equivalent to basic patent US2002182586A1, for which no sequence data was published

Sequence 196686 BP; 53978 A; 42758 C; 43862 G; 55372 T; 0 U; 716 Other;

Query Match 37.5%; Score 149.8; DB 11; Length 196686;

Best Local Similarity 76.3%; Pred. No. 5; 7e-31; Mismatches 58; Indels 2; Gaps 1;

Matches 196; Conservative 1; 81 ATGCTGTATATCCAGCACTTCGGAGGCCAAGTGGCGGATCACTTAGGTCAAGAGA 140

DB 29937 ACGCTTATATCCAGCACTTGGGAGCCGAGGTGAGCGGATCACTTAGGTCAAGAGT 29996
QY 141 TCGAGCAATCTTGGCCCAACATGTTGAACCCCGTCTTTACTATAAATAACAAAATATAGC 200
DB 29997 TCGAGCAATCTTGGCCCAACATGTTGAACCCCGTCTTTACTATAAATAATATAGC 30056
QY 201 TGGGCATGTGTGGCAGACACCTGTAGTCCAGCTACTCAGAGCCGAGATTGACATGAGC 260
DB 30057 TGGGTGTGTGGCAGACACCTGTATCTTACTTACTAGAGGCTGAGGCGAGAGAGA 30116
QY 261 T-GAGATTCGAGAGTGAAGCCGAATATCAGATCAAGAGTGAAGAGTGAAGACCKCCGT 318
DB 30117 TGTATGTAGAGCAAGAGGTGTCATGTGCACTCCAGCTCGGGCAACAGAGTGAAGACTTAA 30176
QY 319 CTCAAAAACAACAACA 335
DB 30177 CTCAAAAAATAAATAA 30193

RESULT 33

ABL65836/C
ID ABL65836 standard; DNA, 5670 BP.

ABL65836;

15-MAY-2002 (first entry)

Lung cancer related gene sequence SEQ ID NO:4173.

Human; cancer; colon; breast; ovary; oesophagus; kidney; thyroid;

stomach; lung; prostate; pancreas; carcinoma; antitumour; cancerous;

cytostatic; gene therapy; antineoplastic; Wilms' tumour; adenocarcinoma;

gene; ds.

Homo sapiens.

WO200194629-A2.

13-DEC-2001.

30-MAY-2001; 2001WO-US010838.

PR 05-JUN-2000; 2000US-0209473P.
PR 05-JUN-2000; 2000US-0209531P.
PR 18-SEP-2000; 2000US-0233133P.
PR 18-SEP-2000; 2000US-0233617P.
PR 20-SEP-2000; 2000US-0234009P.
PR 20-SEP-2000; 2000US-0234034P.
PR 20-SEP-2000; 2000US-0234052P.
PR 22-SEP-2000; 2000US-0234567P.
PR 22-SEP-2000; 2000US-0234923P.
PR 25-SEP-2000; 2000US-0234924P.
PR 25-SEP-2000; 2000US-0234924P.
PR 25-SEP-2000; 2000US-0235077P.
PR 25-SEP-2000; 2000US-0235082P.
PR 25-SEP-2000; 2000US-0235134P.
PR 25-SEP-2000; 2000US-0235280P.
PR 26-SEP-2000; 2000US-0235637P.
PR 26-SEP-2000; 2000US-0235638P.
PR 27-SEP-2000; 2000US-0235711P.
PR 27-SEP-2000; 2000US-0235720P.
PR 27-SEP-2000; 2000US-0235840P.
PR 27-SEP-2000; 2000US-0235863P.
PR 28-SEP-2000; 2000US-0236028P.
PR 28-SEP-2000; 2000US-0236032P.
PR 28-SEP-2000; 2000US-0236032P.
PR 28-SEP-2000; 2000US-0236034P.
PR 28-SEP-2000; 2000US-0236109P.
PR 28-SEP-2000; 2000US-0236111P.
PR 29-SEP-2000; 2000US-0236842P.
PR 29-SEP-2000; 2000US-0236891P.
PR 02-OCT-2000; 2000US-0237172P.

[illegible]

ID	ACA64883	standard; DNA; 5670 BP.
XX	AC	ACA64883;
XX	DT	27-JUN-2003 (first entry)
XX	DE	Human MB-1 gene (CD79a-B cell) DNA corresponding to U05259.
XX	KW	Human; chronic inflammatory joint disease; infection; tumour; antiinflammatory; cytostatic; antiarthritic; antineumatic; immunosuppressive; gene therapy; etiological pathogenicity; ds.
OS	Homo sapiens.	
PX	DN	DEJ0127572-A1.
PX	PD	05-DEC-2002.
PF	30-MAY-2001;	2001DE-01027572.
PR	30-MAY-2001;	2001DE-01027572.
PA	(PATH-) PATHOARRAY GMBH.	
PI	Haenpl T,	Ungethuem U, Blaess S;
DR	WPL; 2003-240797/24.	
PT	Reagents for diagnosis, study and therapy of chronic inflammatory joint and other diseases, comprises any of many specified genes or derived proteins.	
PS	Claim 1; Page; 12pp; German.	
CC	This invention describes a novel reagent for diagnosis, molecular definition and therapy of chronic inflammatory joint diseases, and other inflammatory disorders, infective or tumour diseases in humans. The products of the invention have antiinflammatory, cytostatic, antiarthritic, antineumatic and immunosuppressive activity and can be used for gene therapy. The reagent of the invention and any proteins and antibodies derived from it, are used (i) for analysing tissue and blood samples for medical diagnosis; (ii) for diagnosis and characterisation of chronic joint diseases, on the basis of molecular characteristics, and determining the etiological pathogenicity principle of as yet uncharacterised inflammatory diseases, also monitoring progression and/or treatment of disease, and optimisation of therapy and (iii) for developing treatments for inflammatory diseases, particularly of joints, infections and tumours. ACA64801-ACA64965 represent human polynucleotides used in the method of the invention	
SO	Sequence 5670 BP; 1267 A; 1630 C; 1616 G; 1157 T; 0 U; 0 Other;	
Query Match	37.4%; Score 149.6; DB 8; Length 5670;	
Best Local Similarity	75.0%; Pred. No. 2.2e-31;	
Matches 201; Conservative 1; Mismatches 60; Indels 6; Gaps 1		
QY	83	GCGTGTATCCGAGCACTTCGGGAGGCCAAGSTGGCGCATCCTGAGTCAAGAATC 142
DB	5307	GCGTGTATCCGAGCACTTCGGGAGGCCAAGSTGTGATCCTTGAGTCAAGGTTTC 5244
QY	143	GAGACCATCTCTGGCCAATGATGTGAACCCGCCTTTACTTAATAAAATAAATAAGTCTG 202
DB	5247	AAGACCAATCTGGCCAATGATGTGAACCCGCCTTTACTTAATAAATAAATAAGTCTG 5188
QY	203	GGCATGTGGCAACACCTGTATGTCCTCACTAATCAGAGCCGGAGATTGCACTGAGCTG 262
DB	5187	GGCATGTGGCAACACCTGTATGTCCTCACTAATCAGAGCCGGAGATTGCACTG 5128
QY	263	AGATCGCAGATGAGCCGAATTCACAGATC-----ACAGATGAGCAGATGAGACKC 316
DB	5127	TGAACCCAGAGGAGAGAGTTCGATGAGCACTCCAACCTGGGCAACAGATGAGACTCT 5066
QY	317	GTCTCAAAAACAAACAAAAAACA 344

5067 GTCTCAAAAAAAAAAAAAAAAAA 5040

RESULT	35
AEF74508/c	
ID	AEF74508 standard; DNA; 5670 BP.

AC	AEF74508;
----	-----------

DT	06-APR-2006 (first entry)
----	---------------------------

Human polynucleotide #22.

KM	Diagnosis; gene regulation; gene expression;
KM	post traumatic stress disorder; psychiatric disorder; tranquilizer; gene;
KM	ds.

Homo sapiens.

PN WO2006013561-A2.

PD 09-FEB-2006.

PF 02-AUG-2005; 2005WO-IL000824.

PR 02-AUG-2004; 2004US-0592408P.

(YISS) YISSUM RES DEV CO HEBREW UNIV JERUSALEM

XX

XX

PT New kit comprising 10 and no more than 574 polynucleotides capable of
PT specifically binding at least one specific polynucleotide sequence,
PT useful for determining predisposition of a subject to develop PTSD, or
PT for diagnosing PTSD.

PS	Claim 1; SEQ ID NO 22; 157pp; English.
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The invention relates to a kit for determining predisposition of a subject to developing post-traumatic stress disorder (PTSD) comprising at least 10 and no more than 574 polynucleotides, where each of the polynucleotides is capable of specifically binding at least one specific polynucleotide sequence. The invention also relates to a kit for diagnosing PTSD in a subject, agents for the manufacture of the kits cited comprising the polynucleotides cited, and a microarray comprising at least 10 and no more than 904 oligonucleotides where each of the oligonucleotides is capable of specifically binding at least one specific polynucleotide sequence. The kit comprises each of the polynucleotides selected from an oligonucleotide molecule, a cDNA molecule, a genomic molecule and an RNA molecule. Each of the polynucleotides is at least 10 and no more than 50 nucleic acids in length. Each of the polynucleotides is bound to a solid support. The kit also comprises at least one reagent suitable for detecting hybridization of the polynucleotides and at least one RNA transcript. The kit further comprises packaging materials packaging the at least one reagent and instructions for using the kit in determining predisposition of the subject to developing PTSD, or for diagnosing the disease. The microarray comprises oligonucleotides of at least 10 and no more than 40 nucleic acids in length. The agent is capable of regulating an expression level of at least one gene as a pharmaceutical or for the manufacture of a medicament identified for preventing PTSD. The kit is useful for determining predisposition of a subject to developing PTSD or for diagnosing PTSD. This sequence represents a human polynucleotide of the invention. Note: The sequence data for this patent did not form part of the printed specification, but was obtained in electronic format directly from WIPO at <http://wipo.int/pub/published.pct.sequences>.

AA	Sequence
5670	BP; 1267 A; 1630 C; 1616 G; 1157 T; 0 U; 0 Other;
5671	

Query Match	37.4%	Score 149.6;	DB 15;	Length 5670;
Best Local Similarity	75.0%	Pred. No. 2.2e-31;		
Matches 201; Conservative	1;	Mismatches 60;	Indels 6;	Gaps 1;

QY	8	GCCTGTAATCCACACACTTGGGAGGGCCAAAGTGGGAGATCACCTGAGTCAAGAATC	142
Db	5307	GCTGTATATCCAGACACTTTGGGAGGCCAAGTAGTGAATCACTGAGTCAAGGTTCC	5248
QY	143	GAGACCATCTGTGGCAACATGTGAAAACCCGCTTTTACTAAATAATCAAAAATATAGCTG	202
Db	5247	AAGACCAAGTGTGGCCAAATGTAAGAACCCGCGCTTACTAATAAATACAAAAATTAGCTG	5188
QY	203	GGCATGTGGGACACACCTGTAGTCCAGCACTCAGGAGCCGAGATGTGAGTGAGCTG	262
Db	5187	GGCATGTGGGAGGCGCTGTATATCCAGCTTACTTGGAGGCTGAGGAGGAGAAATCGCT	5128
QY	263	AGATCGCAGATGAGCCGAATTCACAGATC-----ACAGATGAGCAGAGTGAACKCC	316
Db	5127	TGAACCCGAGGAGGAGAGTGTTCAGATGAGCACTCAACCTGGGCAACAGATGAGACTCT	5068
QY	317	GTTCTCAAAAACACACAAAAAACAAAA	344
Db	5067	GTTCTCAAAAACAAAAAACAAAAA	5040

RESULT 36
AAK90749/c
ID AAK90749 standard; DNA; 8205 BP.

AC AAK90749;

DT 05-NOV-2001 (first entry)

Accession	Human digestive system antigen genomic sequence	SEQ ID NO:
DE		4325.

KM Human; digestive system antigen; gene therapy; cancer; appendicitis;
KM ulcerative colitis; infection; Hirschsprung's disease; chronic colitis;
KM digestive system disorder; Meckel's diverticulum; ds.

AA
OS Homo sapiens.

AA WO200155314-A2.
PN

02-AUG-2001

AA 17-JAN-2001; 2001WO-US001324.
PF

PR	14-AUG-2000	2000US-0179065P
PR	31-JAN-2000	2000US-0179065P
PR	04-FEB-2000	2000US-0180628P
PR	24-FEB-2000	2000US-0184664P
PR	02-MAR-2000	2000US-0186350P
PR	16-MAR-2000	2000US-0189874P
PR	17-MAR-2000	2000US-0190076P
PR	18-APR-2000	2000US-0198113P
PR	18-MAY-2000	2000US-0205515P
PR	28-JUN-2000	2000US-0209467P
PR	28-JUN-2000	2000US-0214886P
PR	30-JUN-2000	2000US-0215135P
PR	07-JUL-2000	2000US-0216647P
PR	07-JUL-2000	2000US-0216880P
PR	11-JUL-2000	2000US-0217487P
PR	11-JUL-2000	2000US-0217486P
PR	14-JUL-2000	2000US-0218293P
PR	26-JUL-2000	2000US-0220963P
PR	26-JUL-2000	2000US-0220964P
PR	14-AUG-2000	2000US-0224518P
PR	14-AUG-2000	2000US-0224519P
PR	14-AUG-2000	2000US-0225213P
PR	14-AUG-2000	2000US-0225214P
PR	14-AUG-2000	2000US-0225266P
PR	14-AUG-2000	2000US-0225267P
PR	14-AUG-2000	2000US-0225270P
PR	14-AUG-2000	2000US-0225276P

PR	14-AUG-2000	2000US-02254477
PR	14-AUG-2000	2000US-02255787
PR	14-AUG-2000	2000US-0225758P
PR	14-AUG-2000	2000US-0225759P
PR	18-AUG-2000	2000US-0226279P
PR	22-AUG-2000	2000US-0226661P
PR	22-AUG-2000	2000US-0226688P
PR	22-AUG-2000	2000US-0227182P
PR	23-AUG-2000	2000US-0227700P
PR	30-AUG-2000	2000US-0228924P
PR	01-SEP-2000	2000US-0229287P
PR	01-SEP-2000	2000US-0229343P
PR	01-SEP-2000	2000US-0229354P
PR	01-SEP-2000	2000US-0229345P
PR	05-SEP-2000	2000US-0229509P
PR	05-SEP-2000	2000US-0229513P
PR	06-SEP-2000	2000US-0230437P
PR	06-SEP-2000	2000US-0230438P
PR	08-SEP-2000	2000US-0231242P
PR	08-SEP-2000	2000US-0231243P
PR	08-SEP-2000	2000US-0231244P
PR	08-SEP-2000	2000US-0231413P
PR	08-SEP-2000	2000US-0231414P
PR	08-SEP-2000	2000US-0232080P
PR	08-SEP-2000	2000US-0232081P
PR	12-SEP-2000	2000US-0231968P
PR	14-SEP-2000	2000US-0232397P
PR	14-SEP-2000	2000US-0232398P
PR	14-SEP-2000	2000US-0232399P
PR	14-SEP-2000	2000US-0232400P
PR	14-SEP-2000	2000US-0232401P
PR	14-SEP-2000	2000US-0233063P
PR	14-SEP-2000	2000US-0233064P
PR	14-SEP-2000	2000US-0233065P
PR	21-SEP-2000	2000US-0234233P
PR	21-SEP-2000	2000US-0234274P
PR	25-SEP-2000	2000US-0234997P
PR	25-SEP-2000	2000US-0234998P
PR	26-SEP-2000	2000US-0235484P
PR	27-SEP-2000	2000US-0235834P
PR	27-SEP-2000	2000US-0235835P
PR	29-SEP-2000	2000US-0236327P
PR	29-SEP-2000	2000US-0236367P
PR	29-SEP-2000	2000US-0236368P
PR	29-SEP-2000	2000US-0236369P
PR	29-SEP-2000	2000US-0236370P
PR	02-OCT-2000	2000US-0236802P
PR	02-OCT-2000	2000US-0237037P
PR	02-OCT-2000	2000US-0237038P
PR	02-OCT-2000	2000US-0237039P
PR	02-OCT-2000	2000US-0237040P
PR	13-OCT-2000	2000US-0239935P
PR	13-OCT-2000	2000US-0239937P
PR	20-OCT-2000	2000US-0240960P
PR	20-OCT-2000	2000US-0241212P
PR	20-OCT-2000	2000US-024185P
PR	20-OCT-2000	2000US-024186P
PR	20-OCT-2000	2000US-024187P
PR	20-OCT-2000	2000US-0241808P
PR	20-OCT-2000	2000US-0241809P
PR	20-OCT-2000	2000US-0241826P
PR	08-NOV-2000	2000US-0246533P
PR	08-NOV-2000	2000US-0246534P
PR	08-NOV-2000	2000US-0246535P
PR	08-NOV-2000	2000US-0246536P
PR	08-NOV-2000	2000US-0246527P
PR	08-NOV-2000	2000US-0246528P

Query Match	37.3%	Score 149;	DB 4;	Length 8205;
Best Local Similarity	75.5%	Pred. No. 3.6e-31;		
Matches	200;	Conservative	1;	Mismatches 56; Indels 8; Gaps 1;
QY	81	ATGCTCTGTAATCCCAAGCACTTCGGAGAGCCCAAGTGCGGATCACTCGAGCTCAAGGA	140	
DB	5600	ATGCTCTGTAATCCCAAGCACTTCGGAGAGCCCAAGTGCGGATCACTCGAGCTCAAGGA	140	
QY	141	TCGAGACCATCTGTGGCCCAATGTGGAAACCCGCTTACTTAATAAATATCAAAAATATAGC	200	
DB	5540	TCGAGACCATCTGTGGCCCAATGTGGAAACCCGCTTACTTAATAAATATCAAAAATATAGC	200	
QY	201	TGGGATGTTGGACACACCTGTGATCCCAAGCTACTCAAGAGCGGAGATTGCAAGTGAAC	260	

Db	5480	CGGGTGTGTGTGTGTGTGCCCTTATATCCCACTACTCAGAGGAGGAGGTTTGCAGTAGAC	5421
OY	261	TCAGATCGCAGACGTGAGCCGAATTCACGATTCACAGAGTGAGCAGAGTGCACKCCGTCT	320
Db	5420	CAGAAGCTGC-----GCCATTACACTCCGGCTCGGCAACAAGAGCAAAATCTCATCT	5369
OY	321	CAAAAACACACACAAAAACAAAA	345
Db	5368	CAAAAAAAAAAGAAAAAAAAAGAC	5344
RESULT 37			
AAI62948/c			
ID	AAI62948 standard; DNA; 8205 BP.		
XX	AAI62948;		
AC			
DT	22-OCT-2001 (first entry)		
XX			
DE	Human genomic DNA SEQ ID NO 276.		
XX			
KW	Human; nocotropic; neuroprotective; cytosstatic; dermatological; virucide;		
KW	immunosuppressive; antiinflammatory; anti-HIV; antibacterial; vulnerary;		
KW	antiparkinsonian; antisticking; antianemic; antiarthritic; cancer;		
KW	antirheumatic; hepatotropic; cerebroprotective; antiinflammatory;		
KW	antiallergic; antidiabetic; antitumor; anticonvulsant; antifungal;		
KW	antiparasitic; cardiac; immune disorder; cardiovascular disorder;		
KW	neurological disease; infection; nephrotropic; gene therapy; vaccine; ds.		
XX			
OS	Homo sapiens.		
XX			
PN	WO200155449-A1.		
XX			
PD	02-AUG-2001.		
XX			
PE	17-JAN-2001; 2001WO-US001346.		
XX			
PR	31-JAN-2000;	2000US-0179065P.	
PR	04-FEB-2000;	2000US-0180628P.	
PR	19-MAY-2000;	2000US-0205515P.	
PR	07-JUL-2000;	2000US-0216880P.	
PR	14-JUL-2000;	2000US-0218290P.	
PR	14-AUG-2000;	2000US-0225447P.	
PR	01-SEP-2000;	2000US-0229343P.	
PR	06-SEP-2000;	2000US-0230437P.	
PR	08-SEP-2000;	2000US-0231243P.	
PR	25-SEP-2000;	2000US-0234997P.	
PR	29-SEP-2000;	2000US-0236367P.	
PR	13-OCT-2000;	2000US-0239937P.	
PR	08-NOV-2000;	2000US-0246476P.	
PR	08-NOV-2000;	2000US-0246477P.	
PR	08-NOV-2000;	2000US-0246525P.	
PR	08-NOV-2000;	2000US-0246526P.	
PR	17-NOV-2000;	2000US-0246528P.	
PR	17-NOV-2000;	2000US-0249210P.	
PR	17-NOV-2000;	2000US-0249211P.	
PR	17-NOV-2000;	2000US-0249214P.	
PR	17-NOV-2000;	2000US-0249265P.	
PR	01-DEC-2000;	2000US-0250160P.	
PR	01-DEC-2000;	2000US-0250391P.	
PR	05-DEC-2000;	2000US-0251030P.	
PR	05-DEC-2000;	2000US-0251988P.	
PR	05-DEC-2000;	2000US-0256719P.	
PR	06-DEC-2000;	2000US-0251479P.	
PR	08-DEC-2000;	2000US-0251989P.	
PR	08-DEC-2000;	2000US-0251990P.	
PR	11-DEC-2000;	2000US-0254097P.	
XX			
PA	(HUMA-) HUMAN GENOME SCI INC.		
XX			
FI	Rosen CA, Barash SC, Ruben SM;		
XX			

DR	WP1; 2001-476225/51.
XX	Novel plasma membrane associated proteins useful for diagnosing,
PT	treating, preventing and/or prognosing disorders related to the proteins,
PT	including cancer, immune response and neuronal disorders.
PS	Example 2; SEQ ID NO 276; 532pp + Sequence Listing; English.
XX	
CC	The invention relates to novel genes (AA162752-AA162961) and proteins
CC	(AA162962-AA162963) useful for preventing, treating or ameliorating
CC	medical conditions e.g. by protein or gene therapy. The genes are
CC	isolated from a range of human tissues disclosed in the specification.
CC	The nucleic acids, proteins, antibodies and (ant)agonists are useful in
CC	the diagnosis, treatment and prevention of: (a) cancer, e.g. breast and
CC	ovarian cancer and other cancers of the adrenal gland, bone, bone marrow,
CC	breast, gastrointestinal tract, liver, lung, or urogenital; (b) immune
CC	disorders e.g. Addison's disease, allergies, autoimmune haemolytic
CC	anaemia, autoimmune thyroiditis, diabetes mellitus, Crohn's disease,
CC	multiple sclerosis, rheumatoid arthritis and ulcerative colitis; (c)
CC	cardiovascular disorders such as myocardial ischaemias; (d) wound healing
CC	(e) neurological diseases e.g. cerebral anoxia and epilepsy; and (f)
CC	infectious diseases such as viral, bacterial, fungal and parasitic
CC	infections. Note: The sequence data for this patent did not form part of
CC	the printed specification, but was obtained in electronic format directly
CC	from WIPO at ftp.wipo.int/pub/published_pct_sequences
XX	
SEQ	Sequence 8205 BP; 2216 A; 1646 C; 1763 G; 2580 T; 0 U; 0 Other;
Query Match	37.3%; Score 149; DB 4; Length 8205;
Best Local Similarity	75.5%; Pred. No. 3.6e-31;
Matches	200; Conservative 1; Mismatches 56; Indels 8; Gaps 1;
OY	81 ATGCTTGAATCCCAAGCACTTCGGGAGGCCAAGTGGCGGATACCTGAGTCAAGGA 140
DB	5600 ATGCTTGAATCCCAAGCACTTGAAGGCCAAGCGGGTGAATCACTGAGTCAAGGT 5541
OY	141 TCGAGACCATCTGGCCCAACATGTGTAAACCCCTCTTAAATAAATACAAAATATGCG 200
DB	5540 TCGAGACCATCTGGCCCAACATGTGTAAACCCCTCTTAAATAAATACAAAATATGCG 5481
OY	201 TGGCGATGTGTGGACACACACTGTAGTCCAGCTACTCAGAGCGCGAGATTGCAGTGAAC 260
DB	5480 CGGGTGTGTGTGTGTGGCGCTATATATCCAGCTACTCAGAGCGCGAGATTGCAGTGAAC 5421
OY	261 TGAATGCCAAGTGAAGCCGAATTCACAGATTCACAGAGTGAAGTGAAGTCCCTCT 320
DB	5420 CAAAGCTGTC-----GCCATTACACTCCCGCTGGGCAACAAAGCAAACTCCATCT 5365
OY	321 CAAAACCAACAAACAAAACAAAAA 345
DB	5368 CAAAACCAACAAACAAAACAAAAA 5344
RESULT 38	
ID	ADO79404 standard; DNA; 89900 BP.
XX	ADO79404;
AC	26-AUG-2004 (first entry)
DT	DPF3 region, SEQ ID 3.
XX	
XX	Cytostatic; Gene therapy; breast cancer; human; DGL1; KIAA0783; DPF3;
KW	CENCL1; gene; ds; SNP; single nucleotide polymorphism;
KW	D4; zinc and double PHD fingers, family 3; CERD4; cer-d4; FLJ14079;
KW	2810403B03R1k; Rho family guanine-nucleotide exchange factor;
KW	chromosome 14q24.3-q31.1.
XX	
OS	Homo sapiens.
XX	
PH	Key
FT	variation
FT	160

FT	variation	/*tag= a /standard_name= "Single nucleotide polymorphism" /note= "This SNP is described as a A/C SNP" 6053
FT	variation	/*tag= b /standard_name= "Single nucleotide polymorphism" /note= "This SNP is described as a T/G SNP" 9719
FT	variation	/*tag= c /standard_name= "Single nucleotide polymorphism" /note= "This SNP is described as a A/G SNP" 10481
FT	variation	/*tag= d /standard_name= "Single nucleotide polymorphism" /note= "This SNP is described as a T/C SNP" 10676
FT	variation	/*tag= e /standard_name= "Single nucleotide polymorphism" /note= "This SNP is described as a A/T SNP" 17179
FT	variation	/*tag= f /standard_name= "Single nucleotide polymorphism" /note= "This SNP is described as a C/G SNP" 18561
FT	variation	/*tag= g /standard_name= "Single nucleotide polymorphism" /note= "This SNP is described as a A/T SNP" 18658
FT	variation	/*tag= h /standard_name= "Single nucleotide polymorphism" /note= "This SNP is described as a G/C SNP" 18694
FT	variation	/*tag= i /standard_name= "Single nucleotide polymorphism" /note= "This SNP is described as a A/G SNP" 18858
FT	variation	/*tag= j /standard_name= "Single nucleotide polymorphism" /note= "This SNP is described as a T/C SNP" 24582
FT	variation	/*tag= k /standard_name= "Single nucleotide polymorphism" /note= "This SNP is described as a G/A SNP" 24683
FT	variation	/*tag= l /standard_name= "Single nucleotide polymorphism" /note= "This SNP is described as a G/A SNP" 24767
FT	variation	/*tag= m /standard_name= "Single nucleotide polymorphism" /note= "This SNP is described as a T/C SNP" 27402
FT	variation	/*tag= n /standard_name= "Single nucleotide polymorphism" /note= "This SNP is described as a A/G SNP" 28150
FT	variation	/*tag= o /standard_name= "Single nucleotide polymorphism" /note= "This SNP is described as a T/G SNP" 28494
FT	variation	/*tag= p /standard_name= "Single nucleotide polymorphism" /note= "This SNP is described as a T/C SNP" 32003
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FT	variation	/*tag= w /standard_name= "Single nucleotide polymorphism" /note= "This SNP is described as a G/A SNP" 40033
FT	variation	/*tag= x /standard_name= "Single nucleotide polymorphism" /note= "This SNP is described as a T/G SNP" 40095
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FT	variation	/*tag= z /standard_name= "Single nucleotide polymorphism" /note= "This SNP is described as a A/C SNP" 42799
FT	variation	/*tag= aa /standard_name= "Single nucleotide polymorphism" /note= "This SNP is described as a A/G SNP" 43090
FT	variation	/*tag= ab /standard_name= "Single nucleotide polymorphism" /note= "This SNP is described as a G/A SNP" 46683
FT	variation	/*tag= ac /standard_name= "Single nucleotide polymorphism" /note= "This SNP is described as a A/G SNP" 49774
FT	variation	/*tag= ad /standard_name= "Single nucleotide polymorphism" /note= "This SNP is described as a A/G SNP" 51796
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FT	variation	/*tag= ag /standard_name= "Single nucleotide polymorphism" /note= "This SNP is described as a T/C SNP" 53971
FT	variation	/*tag= ah /standard_name= "Single nucleotide polymorphism" /note= "This SNP is described as a A/C SNP" 55899
FT	variation	/*tag= ai /standard_name= "Single nucleotide polymorphism" /note= "This SNP is described as a T/C SNP" 60682
FT	variation	/*tag= aj /standard_name= "Single nucleotide polymorphism" /note= "This SNP is described as a G/A SNP" 61291
FT	variation	/*tag= ak /standard_name= "Single nucleotide polymorphism" /note= "This SNP is described as a T/C SNP" 72720
FT	variation	/*tag= al /standard_name= "Single nucleotide polymorphism" /note= "This SNP is described as a T/C SNP" 72720

FT	/note= "This SNP is described as a G/A SNP"
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FT	/note= "This SNP is described as a A/G SNP"
FT	variation
FT	89751
FT	/*tag= an
FT	/standard_name= "Single nucleotide polymorphism"
FT	/note= "This SNP is described as a T/A SNP"
XX	
PN	WO2004047514-A2.
PD	10-JUN-2004.
XX	
PE	25-NOV-2003; 2003MO-US037943.
XX	
PR	25-NOV-2002; 2002US--0429136P.
XX	
PR	24-JUL-2003; 2003US-0490234P.
XX	
PA	(SEQU-) SEQUENOM INC.
XX	
PI	Roth RB, Nelson MR, Braun A, Kammerer SM, Reneland R;
XX	
DR	WPI, 2004-441037/41.
XX	
PT	Identifying a subject at risk of breast cancer by detecting the presence
PT	of polymorphic variations in the DLG1, KIAA0783, DPF3 or CENPC1 regions
PT	which are associated with breast cancer in a nucleic acid sample from a
XX	subject.
PS	Claim 24; Fig 3; 227pp; English.
XX	
XX	
Query Match	37.3%; Score 149; DB 12; Length 89900;
Best Local Similarity	76.4%; Pred. No. 7.5e-31;
Matches 207; Conservative	1; Mismatches 61; Indels 2; Gaps 2;
OY	81 ATGCCTGTAAATCCCAAGCACTTGGGAGGCCAAGGTGGCGGATCACTGAAGTCAGAAGA 140
DB	3670 ATGCGTGTAATCCCAAGCACTTGGGAGGCTGAGGTGGGTGATCCCTGAGGTCAGAGT 3722
OY	141 TCAGACCATCCTGCACCACAATGTGTAAACCCCGCTTTACTTAAATAATACAAAATATAGC 200
DB	3730 TCGAACACCAAGCTTGCCCAACATGTGTGACACCCCCTCTCTACTTAAAAATACAAAATATTGAC 3788
OY	201 TGGGCGATGGTGGCACACACTGTAGTCCCAAGTCTACTCAAGAGCCG-GAGATTGCAGTAGG 259
DB	3790 TGGGCGCTGGTGGCAACAGCGCTGTAAATCCCAAGCTACTCAAGAGTCCCTTGAACCTCAGAAAG 3843
OY	260 CTGAGATGCGACAGATGAGGCGCAAATACACGATCACAGAGTAGG-CAGAGTGGAGACKCGT 318
DB	3850 TTGTGTGTGAGTTGGAGATTGTGCCACTGCACCTCAGCGCTGGCGAACAGGAGTGAAGTCTGT 3905
OY	319 CTCAAAAACAACACAAAAAAACMAAAAAACC 349
DB	3910 CTAAAAAAAAAAAAAAAAAAAAAAAAAAAC 3940
RESULT 39	
AEP92655/c	
ID	AEP92655 standard; cDNA; 174318 BP.
XX	
AC	AEP92655;
XX	
DT	20-APR-2006 (first entry)
DE	Human Ras effector protein Rin3, cDNA.
XX	
XX	Diagnostics; ss; gene; neurodegenerative disease; Alzheimers disease;

KM	dementia; cognitive disorder; Parkinsons disease; microarray;
XX	neuroprotective; nootropic; antiparkinsonian; screening; prognosis.
OS	Homo sapiens.
XX	
PN	MO2006020269-A2.
XX	
PD	23-FEB-2006.
XX	
XX	19-JUL-2005; 2005MO-US025491.
PF	19-JUL-2004; 2004US-0589318P.
PR	
XX	(UVRP) UNIV ROCHESTER.
PA	
XX	Coleman PD, Federoff HJ, Maguire-Zeiss K, Myhre TR, Kurian RM;
PI	Cox C, Marshall F;
XX	
DR	WPI; 2006-184393/19.
XX	
PT	Use of biomarkers for neurodegenerative disease for, e.g. diagnosing
PT	neurodegenerative disease, screening therapeutic agent for treating
PT	neurodegenerative disease, or monitoring neurodegenerative disease
PT	progression.
XX	
PS	Example 1; SEQ ID NO 82; 552pp; English.
XX	
CC	The invention relates to using a biomarker for a neurodegenerative
CC	disease for diagnosing a neurodegenerative disease, screening a
CC	therapeutic agent for treating a neurodegenerative disease, monitoring a
CC	neurodegenerative disease progression, monitoring a response to a
CC	neurodegenerative disease treatment, identifying a risk for a
CC	neurodegenerative disease, and differentially diagnosing a
CC	neurodegenerative disease in a test subject. Also included are diagnosing
CC	a neurodegenerative disease in a subject, screening for a therapeutic
CC	agent for the treatment of a neurodegenerative disease, monitoring a
CC	neurodegenerative disease progression in a subject, monitoring a response
CC	to a neurodegenerative disease treatment in a subject, identifying a risk
CC	for a neurodegenerative disease in a test subject, differentially
CC	diagnosing a neurodegenerative disease in a test subject, a solid support
CC	(comprising one or more biomarkers, where the biomarker is one or more
CC	proteins comprising HSP60, Dihydropyrimidine dehydrogenase, ER-60
CC	protease, Glucose-6-phosphate dehydrogenase, Atp-synthase beta chain,
CC	Annexin I, 14-3-3 epsilon, Prohibitin, Phosphoglycerate mutase 1, lactoy
CC	Apolipoprotein AI, Superoxide dismutase, RNA-binding protein regulator
CC	subunit, Chain A thioester oxidase B, RAS-related protein RAP1B,
CC	Tumor rejection antigen, Haptoglobin, Fibrin beta, or its combinations)
CC	and a solid support comprising one or more biomarkers (where the
CC	biomarker is one or more transcripts comprising cyclin D1, cyclin B,
CC	cyclin G1, weel, hHR23, CDC25B, GSK3 beta, protein kinase C alpha, C5, C1
CC	inhibitor, IL-11R, IL-8, LIF, TNF-alpha, IL-10R, Alpha-1
CC	antichymotrypsin, HSP 27, HSP 90, crystalline, GAPDH, ferritin H,
CC	ferritin L, cox 1, cox 2, transferrin, or its combinations). The
CC	biomarkers for the neurodegenerative disease are useful for diagnosing a
CC	neurodegenerative disease, screening a therapeutic agent for treating a
CC	neurodegenerative disease, monitoring a neurodegenerative disease
CC	progression monitoring a response to a neurodegenerative disease
CC	treatment, identifying a risk for a neurodegenerative disease, and
CC	differentially diagnosing a neurodegenerative disease, e.g. Alzheimer's
CC	disease and Parkinson's disease, in a test subject. The present sequence
CC	is a cDNA for a human biomarker, used in a microarray in the method of
CC	the invention. NOTE: The specification describes AEF92574-AEF92813 (table
CC	4) as cDNA sequences yet some are protein sequences and some genomic DNA.
XX	
SQ	Sequence 174318 BP; 43826 A; 41095 C; 42605 G; 46792 T; 0 U; 0 Other;
XX	
Query Match	37.3%; Score 149; DB 15; Length 174318;
Best Local Similarity	73.5%; Pred. No. 9, 2E-11;
Matches 202; Conservative	1; Mismatches 71; Indels 1; Gaps 1;
0Y	84 CCTGTAATCCAGCACTTCGGGAGGCCAGGTGGCGGATCACTGAGTCAAGAGATCG 143
DB	CCCTTAATGCCAGCACTTTGGAAGGCCGAGTGGAGGATCACTGAAGGTCAAGAGATCG 829020

OY	144	AGACCATCTGCGCAACATGCTGAAACCCCGTCTTACTAAATAATACAAATAATAGCTG	203	
DB	82901	AGACCAGCCTGGCCAAATGTAAGAAACCGTCTCTGCTAAATAATACAAATAATAGCCAG	82842	
OY	204	GCATGCTGGCAACACCTTACTGCTCCAGCTACTCAGAGACCGGAGATTGCATGAGCTGA	263	
DB	82841	GGTGCTGGCAACGCTGTAGTCCAGCTACTTGGAGAGCTGACGACAGAAATGCTT	82782	
OY	264	GATCGCAGAGTAGCGCGAATATCAGATCAAGAGTAG- CAGAGTAGACKCCGCTCTCA	322	
DB	82781	GAACCCAGAGAGGTGAGAGGTTCAGTGAAGCTGAGATTGCCCACTGCAGAGACTGTCTCA	82722	
OY	323	AAACACACACAAAAAACAACATPAGACA	357	
DB	82721	CCAAAAAAGAAAAAGAAAAAGAAAAA	82687	
RESULT	40			
ID	ADX98573.			
AD	ADX98573	standard; DNA; 285300 BP.		
AG	ADX98573;			
XX				
XX	05-MAY-2005	(first entry)		
DB		Human D4, zinc and double PHD fingers, family 3 (DPF3) genomic DNA.		
XX				
XX				
KY		SNP detection; breast tumor; endocrine disease;		
KY		gynecology and obetetics; neoplasm; cytostatic; metastasis;		
KY		gene therapy; RNA interference; chromosome 14; ds; SNP;		
KY		single nucleotide polymorphism;		
KY		D4, zinc and double PHD fingers, family 3; DPF3;		
XX		guanine-nucleotide exchange factor.		
XX				
OS		Homo sapiens.		
PH	Key	Location/Qualifiers		
FT	variation	207		
FT		/tag= a		
FT		/standard_name= "Single nucleotide polymorphism (SNP) "		
FT	variation	486		
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FT		/standard_name= "Single nucleotide polymorphism (SNP) "		
FT	variation	1745		
FT		/tag= c		
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FT	variation	1922		
FT		/tag= d		
FT		/standard_name= "Single nucleotide polymorphism (SNP) "		
FT	variation	2190		
FT		/tag= e		
FT		/standard_name= "Single nucleotide polymorphism (SNP) "		
FT	variation	2590		
FT		/tag= f		
FT		/standard_name= "Single nucleotide polymorphism (SNP) "		
FT	variation	2637		
FT		/tag= g		
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FT	variation	2804		
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FT	variation	2806		
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FT	variation	2895		
FT		/tag= j		
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FT	variation	3185		
FT		/tag= l		
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FT      variation
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FT      variation
FT      /tag= am
FT      /standard_name= "Single nucleotide polymorphism (SNP)"
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FT      /tag= an
FT      /standard_name= "Single nucleotide polymorphism (SNP)"
FT      14507
FT      variation
FT      /tag= ao
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FT      /standard_name= "Single nucleotide polymorphism (SNP)"
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FT      /tag= ba
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FT      21905
FT      variation
FT      /tag= bb
FT      /standard_name= "Single nucleotide polymorphism (SNP)"
FT      22252
FT      variation
FT      /tag= bc
FT      /standard_name= "Single nucleotide polymorphism (SNP)"
FT      22941
FT      variation
FT      /tag= bd
FT      /standard_name= "Single nucleotide polymorphism (SNP)"
FT      23542
FT      variation
FT      /tag= be
FT      /standard_name= "Single nucleotide polymorphism (SNP)"
FT      24677
FT      variation
FT      /tag= bf
FT      /standard_name= "Single nucleotide polymorphism (SNP)"
FT      25009
FT      variation
FT      /tag= bg
FT      /standard_name= "Single nucleotide polymorphism (SNP)"
FT      25618
FT      variation
FT      /tag= bh
FT      /standard_name= "Single nucleotide polymorphism (SNP)"
FT      26082
FT      variation
FT      /tag= bi
FT      /standard_name= "Single nucleotide polymorphism (SNP)"
FT      26136

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FT      /tag= bj
FT      Query Match 37.3%; Score 149; DB 14; Length 285300;
FT      Best Local Similarity 76.4%; Pred. No. 1.1e-30;
FT      Matches 207; Conservative 1; Mismatches 61; Indels 2; Gaps 2;
Oy      81 ATGCGTAAATCCAGCACTTCGGAGGCGCAAGGTGGGCGGATCACCTGAGTCAAGGA 140
Dy      164620 ATGCGTAAATCCAGCACTTCGGAGGCGTGAAGTGGGATGCGCTGAGGTCAAGGA 164679
Oy      141 TCGAGACCATCTGGCCCAACATGCTGAACCCCGCTTTTACTAAATACAAAAATAGC 200
Dy      164680 TCGAGACCATCTGGCCCAACATGCTGAACCCCGCTTTTACTAAATACAAAAATAGC 164739
Oy      201 TGGGCATGTGGGCAACACCGTGTGTCCTCCAGCTACTGAGAGCGG-GAGATTGAGTGA 259
Dy      164740 TGGGCATGTGGGCAACACCGTGTGTCCTCCAGCTACTGAGAGCGG-GAGATTGAGTGA 164799
Oy      260 CTGAGATCGCAGAGTGAAGCCGAATTCACAGATCAAGAGTGAAG-CAGAGTGAAGCKCCGT 318
Dy      164800 TTGTGTGAGTGTGGGATTGTGCCACTGCACTCCAGCCCTGGGCGACAGGGTGAAGTCTGT 164859
Oy      319 CTCAAAAACACACAAACAAAAACAAAAAACC 349
Dy      164860 CTCAAAAACACACAAACAAAAACAAAAAACC 164890

RESULT 41
ABK83568/c
ID ABK83568 standard; DNA; 201143 BP.
XX
AC ABK83568;
XX
Dy      29-AUG-2002 (first entry)
XX
Dy      Human DNA differentially expressed in granulocytic cells #139.
XX
DE Human; ds; granulocytic cell; DNA chip; bacterial infection;
XX
KW viral infection; parasitic infection; protozoal infection;
XX
KW fungal infection; sterile inflammatory disease; psoriasis;
XX
KW rheumatoid arthritis; glomerulonephritis; asthma; thrombosis;
XX
KW cardiac reperfusion injury; renal reperfusion injury; AIDS;
XX
KW adult respiratory distress syndrome; inflammatory bowel disease;
XX
KW Crohn's disease; ulcerative colitis; periodontal disease;
XX
KW granulocyte activation; chronic inflammation; allergy.
OS Homo sapiens.
XX
PN WO200228999-A2.
XX
PD 11-APR-2002.
XX
PF 03-OCT-2001; 2001WO-US030821.
XX
PR 03-OCT-2000; 2000US-0237189P.
XX
PA (GENE-) GENE LOGIC INC.
XX
PI Beazer-Barclay Y, Weissman SM, Yamaga S, Vockley J;
XX
DR WPI; 2002-435328/46.
XX
PT Detecting granulocyte activation by detecting differential expression of
XX
PT genes associated with granulocyte activation, which serves as diagnostic
XX
PT markers that is useful for monitoring disease states and drug toxicity.
XX
PS Claim 1; SEQ ID NO 139; 114bp; English.
CC The invention relates to detecting (M1) granulocyte (GC) activation
CC (GCA), by detecting the level of expression of gene(s) (Gs) identified by
CC DNA chip analysis as given in the specification, and comparing the
CC expression level to an expression level in an unactivated GC, where
CC differential expression of Gs is indicative of GCA. Also included are

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modulating (M2) GA by contacting GC with an agent that alters the expression of at least one gene in Gs; (2) screening (M3) for an agent capable of modulating GCA or an inflammation (especially chronic) in a tissue, an allergic response in a subject, exposure of a subject to a pathogen or sterile inflammatory disease using the gene expression profile; (3) detecting (M4) an inflammation (especially chronic) in a tissue, an allergic response in a subject, exposure of a subject to a pathogen or sterile inflammatory disease, by detecting the level of expression in a sample of the tissue of gene(s) from Gs, where the level of expression of the gene is indicative of inflammation; (4) treating (M5) an inflammation (especially chronic) or in a tissue, an allergic response in a subject, exposure of a subject to a pathogen or sterile inflammatory disease, by contacting a tissue having inflammation with an agent that modulates the expression of gene(s) from Gs in the tissue. M1 is useful for detecting GCA; M2 is useful for modulating Gs; M3 is useful for screening an agent capable of modulating GCA preferably in an inflammation in a tissue; M4 is useful for detecting an inflammation (especially chronic) in a tissue, an allergic response in a subject, exposure of a subject to a pathogen or sterile inflammatory disease (e.g. psoriasis, rheumatoid arthritis, glomerulonephritis, asthma, thrombosis, cardiac reperfusion injury, renal reperfusion injury, ARDS, adult respiratory distress syndrome, inflammatory bowel disease, Crohn's disease, ulcerative colitis, peritonitis, inflammatory disease, also bacterial infection, viral infection, parasitic infection, protozoal infection, fungal infection and M5 is useful for treating one of the above conditions. The present sequence represents a gene differentially expressed in granulocytes. Note: The sequence data for this patent did not form part of the printed specification, but was obtained in electronic format directly from WIPO at http://wipo.int/pub/published_pat_sequences

Query Match	37.2%	Score 148.8;	DB 6;	Length 201143;
Best Local Similarity	70.7%	Pred. No. 1.1e-30;		
Matches 229; Conservative	1;	Mismatches 83;	Indels 11;	Gaps 2

QY	3	AAATATTAATGA	GACATTTGTCAGGCGAGGCATGACATCGGCTGAAATGCTGTAAATCCGAC	96	
Db	3229	ACATTTTAACAATATTAATTAATTCAGGCGAGGCATG---	GTGGCTCATGCTGTAAATCCGACG	317	
QY	97	CACCTTCGGAGGCCAAGGTGGGCGGATCA	CCTGAGGTCAAGAGATCGAGACCATCTGGC	156	
Db	3173	CACCTTTTGAGAGCTGAGACTGGCAGATCA	CTTGAGGTCAAGAGATTGAGACCAAGCTTGGC	311	
QY	157	CAACATGTGTAA	CCCCGCTCTTTACTTAAATAACAAAATAATAGTCGGCATGTGGCA	216	
Db	3113	CAACATGTGTAA	CCCCCATCTCTTACTTAAATAATCAAAAATTGACAGAGCATGTGGCAGG	305	
QY	217	CACCTGTATGCCAGCTACTGAG	-----GCCGGAGATTGCAGTGAAGCTGAGATCGC	269	
Db	3053	CACCTGTATGCCAGCTACTGAGAGGCTGAGGCGAGGAGAAATCGCTTGAA	CCCCCTGGAGGC	299	
QY	270	AGAGTGAGCCGAATC	CACAGATCA	CAGAGTGAGCGAGTGAAGACACCGTCTTAAAAACAA	329
Db	2993	GGAGTTTTCAGGCACTGAGTGGCAT	TGGCGACAGAGCAAACTCTGTCAAAAAA	293	
QY	330	CAACAAAAACAAAAAACATAA	353		
Db	2933	AAATTTGCTGAGTAAATGAACCAAAA	2910		

XX	RESULT 42
AD	ADA02774/c
XX	ID
XX	ADA02774 standard; DNA; 44075 BP.
AD	ADA02774;
XX	
XX	06-NOV-2003 (first entry)
XX	
XX	Human ARHGEF1 carcinoma associated gene, SFO ID NO:1292.
XX	
XX	Human; carcinoma associated; oncogene; carcinoma; cancer; breast;

KW prostate; lymphoma; leukaemia; cytostatic; gene therapy; drug screening,
 KW gene; ds.
 XX
 XX Homo sapiens.
 OS
 XX WO2003057146-A2.
 PN
 XX
 XX
 PD 17-JUL-2003.
 XX
 XX
 PP 26-DEC-2002; 2002WO-US041414.
 XX
 XX 26-DEC-2001; 2001US-00035832.
 PR
 XX
 PA (SAGR-) SAGRES DISCOVERY.
 XX
 XX
 PI Morris DW;
 XX
 XX WPI; 2003-587068/55.
 DR
 XX
 XX
 PT New recombinant nucleic acid encoding carcinoma associated protein,
 useful for preparing compositions for treating carcinomas.

Claim 1; SEQ ID NO 1292; 245pp; English.

CC The invention relates to recombinant carcinoma associated (CA) nucleic
CC acid sequences from mouse and human (ADA01482-ADA03094), and to
CC recombinant carcinoma associated proteins (CAP) encoded by them. The
CC invention also encompasses expression vectors and host cells comprising a
CC CA nucleic acid, a polypeptide (especially an antibody) that specifically
CC binds to the protein, and a bioclip comprising CA nucleic acid or
CC fragments thereof. The sequences of the invention were identified using
CC oncogenic retroviruses, which insert into the genome of the host organism
CC at random. Many of these do not carry transduced host oncogenes or
CC pathogenic trans-acting viral genes, meaning that cancer incidence is a
CC direct consequence of the effects of proviral integration into host
CC protooncogenes. The CA nucleic acid sequences can be used to diagnose
CC carcinoma (especially breast cancer, prostate cancer, lymphoma or
CC leukemia) or a propensity to carcinoma by determination of the sequence
CC of a CA gene, or by determination of CA gene expression in particular
CC tissues. CA nucleic acids, proteins and antibodies are also useful as
CC therapeutic agents and in screening and evaluating drug candidates. The
CC present sequence represents a specifically claimed human CA nucleic acid
CC sequence of the invention. Note: The complete sequence data for this
CC patent did not form part of the printed specification, but was obtained
CC in electronic format directly from WIPO at
CC ftp.wipo.int/pub/published_pat_sequences.

SQ Sequence 44075 BP; 8314 A; 11402 C; 11137 G; 8371 T; 0 U; 4851 Other;

Query Match	37.2%;	Score 148.6;	DB 9;	Length 44075;
Best Local Similarity	74.9%;	Pred. No. 7.7e-31;		
Matches 200;	Conservative 1;	Mismatches 60;	Indels 6;	Gaps 1

[illegible]

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RESULT 43
ADB72512/C
ID ADB72512 standard; DNA; 44075 BP.
XX
XX
AC ADB72512;
XX
XX 04-DEC-2003 (first entry)
DT
XX
DE Human ARHGEF1 gene.
XX
XX human; ds; cytosolic; gene therapy; vaccine; carcinoma; lymphomas;
XX cancer; neoplasm; adenocarcinoma; sarcoma; gene.
XX
XX Homo sapiens.
XX
XX MO2003008583-A2.
XX
XX 30-JAN-2003.
XX
XX 26-DEC-2001; 2001WO-US051291.
XX
XX 02-MAR-2001; 2001US-00798586.
XX 23-OCT-2001; 2001US-00004113.
XX 08-NOV-2001; 2001US-00052482.
XX 30-NOV-2001; 2001US-00997722.
XX 20-DEC-2001; 2001US-00034650.
XX
XX (SAGR-) SAGRES DISCOVERY.
XX
XX Morris DW, Engelhard EK;
XX
XX WPI; 2003-239337/23.
XX
XX New recombinant nucleic acid, useful for treating carcinomas, lymphomas,
XX cancers, neoplasm, adenocarcinoma, or sarcomas.
XX
XX Claim 1; SEQ ID NO 340; 2304pp; English.
XX
XX The invention relates to a novel recombinant nucleic acid comprising a
XX nucleotide sequence selected from any of the 660 sequences fully defined
XX in the specification. A polynucleotide of the invention has cytosolic
XX activity, and may have a use in gene therapy, or in a vaccine. The
XX recombinant nucleic acids and polypeptides are useful for treating
XX carcinomas, e.g. lymphomas, cancers, neoplasm, adenocarcinoma, and
XX sarcomas. The present sequence represents a human gene of the invention.
XX
XX Sequence 44075 BP; 8314 A; 11402 C; 11137 G; 8371 T; 0 U; 4851 Other;
SQ
Query Match 37.2%; Score 148.6; DB 10; Length 44075;
Best Local Similarity 74.9%; Pred. No. 7.7e-31;
Matches 200; Conservative 1; Mismatches 60; Indels 6; Gaps 1;
QY 83 GCGTGTATCCAGCACTTCGGAGGCCAAGGTGGCGGATCAGTCAAGATC 142
DB 8665 GCGTGTATCCAGCACTTCGGAGGCCAAGGTGGCGGATCAGTCAAGATC 8606
QY 143 GAGACATCTGCGCAACATGTGAAACCCGCTTTTACTAAAAATACAAAAATAGCTG 202
DB 8605 AAGACCAATCTGCGCAACATGTGAAACCCGCTTTTACTAAAAATACAAAAATAGCTG 8546
QY 203 GGCATGTGGGACACACCTGTGATCCAGCTACTCAGGAGCGGAGATTGACAGCTG 262
DB 8545 GGCATGTGGGACAGGCGCTGTGATCCAGCTACTCAGGAGCGGAGATTGACAGCTG 8486
QY 263 AGATCGCAGAGTGAGCCGAATTCACAGATC-----ACAGAGTGACAGAGTGAGACKCC 316
DB 8485 TGAACCCAGAGGACAGAGGTTGCAGTGACACTCCAACTGGGCAACAGAGTGAGACTCT 8426
QY 317 GTCTCAAAAACACACACAAAAACAAA 343
DB 8425 GTCTCAAAAACAAAAA 8399
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```
RESULT 44
ADC85254/C
ID ADC85254 standard; DNA; 44075 BP.
XX
XX
AC ADC85254;
XX
XX 01-JAN-2004 (first entry)
DT
XX
DE Human Arhgef1 genomic sequence.
XX
XX Cytosolic; gene therapy; vaccine; cancer; carcinoma-associated gene; CA;
XX secreted; transmembrane; intracellular; ds.
XX
XX Homo sapiens.
XX
XX MO2003045230-A2.
XX
XX 05-JUN-2003.
XX
XX 02-DEC-2002; 2002WO-US038582.
XX
XX 30-NOV-2001; 2001US-00997722.
XX
XX (SAGR-) SAGRES DISCOVERY.
XX
XX Morris DW, Engelhard EK;
XX
XX WPI; 2003-513603/48.
XX
XX New recombinant nucleic acid comprising a nucleotide sequence of any of
XX the carcinoma-associated (CA) genes, useful for screening for drug
XX candidates for diagnosing or treating carcinomas.
XX
XX Claim 1; SEQ ID NO 40; 983pp; English.
XX
XX The invention relates to a recombinant nucleic acid comprising a
XX nucleotide sequence selected from any of the fully defined carcinoma-
XX associated (CA) genes from the 50 tables given in the specification. The
XX CA proteins are secreted, transmembrane or intracellular proteins. The
XX recombinant nucleic acids are useful for screening for drug candidates
XX for diagnosing or treating carcinomas. Sequences given in ADC85215-
XX ADC85514 represent CA genes of the invention.
XX
XX Sequence 44075 BP; 8314 A; 11401 C; 11138 G; 8371 T; 0 U; 4851 Other;
SQ
Query Match 37.2%; Score 148.6; DB 10; Length 44075;
Best Local Similarity 74.9%; Pred. No. 7.7e-31;
Matches 200; Conservative 1; Mismatches 60; Indels 6; Gaps 1;
QY 83 GCGTGTATCCAGCACTTCGGAGGCCAAGGTGGCGGATCAGTCAAGATC 142
DB 8665 GCGTGTATCCAGCACTTCGGAGGCCAAGGTGGCGGATCAGTCAAGATC 8606
QY 143 GAGACATCTGCGCAACATGTGAAACCCGCTTTTACTAAAAATACAAAAATAGCTG 202
DB 8605 AAGACCAATCTGCGCAACATGTGAAACCCGCTTTTACTAAAAATACAAAAATAGCTG 8546
QY 203 GGCATGTGGGACACACCTGTGATCCAGCTACTCAGGAGCGGAGATTGACAGCTG 262
DB 8545 GGCATGTGGGACAGGCGCTGTGATCCAGCTACTTGGAGGCTGAGGCAAGAAATCGCT 8486
QY 263 AGATCGCAGAGTGAGCCGAATTCACAGATC-----ACAGAGTGACAGAGTGAGACKCC 316
DB 8485 TGAACCCAGAGGACAGAGGTTGCAGTGACACTCCAACTGGGCAACAGAGTGAGACTCT 8426
QY 317 GTCTCAAAAACACACACAAAAACAAA 343
DB 8425 GTCTCAAAAACAAAAA 8399
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```
RESULT 45
ADM74369/C
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ID ADM74369 standard; DNA; 44075 BP.
 AG ADM74369;
 XX
 XX 01-JUL-2004 (first entry)
 XX
 XX Human carcinoma associated (CA) nucleic acid #19.
 DB
 XX Human; carcinoma associated nucleic acid; CA nucleic acid; gene; de;
 KM carcinoma associated protein; CAP; carcinoma; leukemia; lymphoma;
 KM cytosolic.
 XX
 XX Homo sapiens.
 OS
 XX US2004072154-A1.
 XX
 XX 15-APR-2004.
 XX
 XX 30-NOV-2001; 2001US-00997722.
 XX
 XX 22-DEC-2000; 2000US-00747377.
 XX
 XX 02-MAR-2001; 2001US-00798586.
 XX
 XX (MORRIS D W.
 PA (ENGE/ ENGELHARD E K.
 XX
 XX Morris DW, Engelhard EK;
 PI
 XX WPI; 2004-328562/30.
 XX
 XX New carcinoma associated gene or protein, useful for preparing a
 PT composition for diagnosing or treating carcinoma e.g., leukemia or
 PT lymphoma.
 XX
 XX Claim 1; SEQ ID NO 40; 29pp; English.
 XX
 XX The invention relates to new recombinant nucleic acids. The invention
 CC also relates to a host cell comprising a recombinant nucleic acid or
 CC expression vector, an expression vector comprising a recombinant nucleic
 CC acid, a recombinant protein, a method of screening for drug candidates, a
 CC method of screening for a bioactive agent capable of binding to a
 CC carcinoma associated protein (CAP) encoded by a nucleotide sequence, a
 CC method of screening for a bioactive agent capable of modulating the
 CC activity of a CAP, a method of evaluating the effect of a candidate
 CC carcinoma drug, a method of diagnosing carcinoma, a method for inhibiting
 CC the activity of a CAP, a method of treating carcinomas, a method of
 CC neutralizing the effect of a CAP and a method of diagnosing carcinoma or
 CC propensity to carcinoma. A method of evaluating the effect of a candidate
 CC carcinoma drug comprises administering the drug to a patient, removing a
 CC cell sample from the patient and determining alterations in the
 CC expression or activation of a gene comprising the nucleotide sequence. A
 CC method of diagnosing carcinoma comprises determining the expression of
 CC one or more genes comprising the nucleic acid sequence in a first tissue
 CC type of a first individual and comparing the expression of the gene from
 CC a second normal tissue type from the first individual or a second
 CC unaffected individual, where a difference in the expression indicates
 CC that the first individual has carcinoma. A method of inhibiting the
 CC activity of a CAP comprises binding an inhibitor to the CAP. Treating
 CC carcinomas comprises administering to a patient an inhibitor of CAP.
 CC Neutralizing the effect of a CAP comprises contacting an agent specific
 CC for the CAP. The polypeptide specifically binds to the protein encoded by
 CC the nucleic acid. It comprises an antibody that specifically binds to the
 CC protein encoded by the nucleic acid. The nucleic acids are useful for
 CC preparing a composition for diagnosing or treating carcinoma e.g.,
 CC leukemia or lymphoma. This sequence represents a human carcinoma
 CC associated (CA) nucleic acid of the invention. Note: The sequence data
 CC for this patent did not form part of the printed specification but was
 CC obtained in electronic format directly from USPTO at
 CC Seqdata.uspto.gov/sequence.html.
 CC
 XX Sequence 44075 BP; 8314 A; 11402 C; 11137 G; 8371 T; 0 U; 4851 Other;
 SQ Query Match 37.2%; Score 148.6; DB 12; Length 44075;

Best Local Similarity 74.9%; Pred. No. 7.7e-31;
 Matches 200; Conservative 1; Mismatches 60; Indels 6; Gaps 1;
 QY 83 GCTGTATATCCAGACTTGGAGGCGCAAGGTGGCGGATCACCTGAGGTCAAGATC 142
 DB 8665 GCTGTATATCCAGACTTGGAGGCGCAAGGTGGAGTGGATCACCTGAGGTCAAGATC 8606
 QY 143 GAGACATCTGGCCAAATGTTGAAACCCGCTTTTCTTAAATAACAAAAATAGCTG 202
 DB 8605 AAGACCACTCTGGCCAAATGTTGAAACCCGCTTTTCTTAAATAACAAAAATAGCTG 8546
 QY 203 GGCAATGGTGACACACCTGTAGTCCGACTACTCAGAGCGCGAGATTGCACTGAGCTG 262
 DB 8545 GGCAATGGTGACAGGCGCTGTATATCCAGCTACTTGGAGGCTGAGGCAAGATTCGCT 8486
 QY 263 AGATGCGAGAGTGAAGCCGAAATCAAGATC-----ACAGAGTGAGCAGAGTGAGACKCC 316
 DB 8485 TGAACCCAGAGAGGCGAGAGTGTGACAGTGAACCTCAACCTGGGCAACAGAGTGAAGACTCT 8426
 QY 317 GTCTCAAAAAACAAACAAAAACAAA 343
 DB 8425 GTCTCAAAAAACAAAAACAAAAA 8399
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 ACN45134
 ID ACN45134 standard; DNA; 49745 BP.
 AC
 AC ACN45134;
 XX
 XX 18-NOV-2004 (first entry)
 DT
 XX Human genomic sequence hCG31443.
 DE
 XX Cytosolic; carcinoma; lymphoma; cancer; human; gene; ss.
 XX
 XX Homo sapiens.
 XX
 XX W02003073826-A2.
 XX
 XX 12-SEP-2003.
 XX
 XX 28-FEB-2003; 2003WO-US006235.
 XX
 XX 01-MAR-2002; 2002US-00087192.
 XX
 XX (SAGRES-) SAGRES DISCOVERY.
 PA
 XX Morris DW;
 PI
 XX WPI; 2003-328604/31.
 XX
 XX Recombinant nucleic acid useful for diagnosis and treatment of carcinoma
 PT comprises a nucleotide sequence.
 PT
 XX Claim 1; SEQ ID NO 1930; 0pp; English.
 XX
 XX The present invention relates to novel DNA and protein sequences which
 CC are associated with carcinomas. The sequences are useful for: (i) for
 CC screening drug candidates; (ii) for screening of bioactive agent capable
 CC of binding to Carcinoma Associated Protein (CAP); (iii) for screening of
 CC a bioactive agent capable of modulating the activity of CAP; (iv) for
 CC evaluating the effect of a candidate carcinoma drug; (v) for diagnosing
 CC carcinoma; (vi) for inhibiting the activity of CAP; (vii) for treating
 CC carcinoma; (viii) for neutralizing the effect of CAP; (ix) as a biochip;
 CC (x) for diagnosing carcinoma or a propensity to carcinoma; and (xi) for
 CC determining Carcinoma Associated (CA) gene copy number. In addition, the
 CC CA genes are useful as DNA vaccines and the CAP are useful as markers of
 CC carcinoma including lymphoma. The present sequence is one such CA coding
 CC sequence. Note: This patent is an equivalent to basic patent
 CC US2002182586A1, for which no sequence data was published
 XX
 XX Sequence 49745 BP; 10151 A; 14745 C; 14496 G; 10151 T; 0 U; 202 Other;
 SQ

ADH76849;
 22-APR-2004 (first entry)
 Melanin-concentrating hormone receptor 1 locus clone.
 melanin-concentrating hormone receptor 1; MCHRI; SNP;
 single nucleotide polymorphism; anorectic; gene therapy; obesity; gene;
 ds.
 Homo sapiens.
 Synthetic.
 Key
 variation
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 /tag= p
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 /replace(103156,C)
 /tag= q
 /standard_name= "Single nucleotide polymorphism"
 /replace(103270,A)
 /tag= r
 /standard_name= "Single nucleotide polymorphism"

05-JUN-2003; 2003MO-EP005917.
 05-JUN-2002; 2002EP-00012569.
 (UYPH-) UNIV PHILIPPS MARBURG.
 Platzer M, Platzer C, Gudermann T, Hebebrand J, Hinney A;
 Reichwald K;
 WPI; 2004-062377/06.
 New diagnostic composition, useful for diagnosing obesity related to the presence of a molecular variant of the MCHRI gene or a susceptibility to the disorder.
 Claim 1; Page; 76pp; English.
 The invention relates to a novel diagnostic polymnucleotide composition. The polymnucleotide composition comprises: a sequence encoding a polypeptide with defined sequences given in the specification; a sequence capable of hybridizing to a melanin-concentrating hormone receptor 1 (MCHRI) gene; a polymnucleotide encoding an MCHRI polypeptide; or a sequence comprising one or more of the nucleotide exchanges (SNP's) given in the specification and at least 8 bases of surrounding sequence of the MCHRI gene. The composition has anorectic activity. The polymnucleotide composition may be used in gene therapy to treat the disorders of the invention. The composition is useful for diagnosing obesity related to the presence of a molecular variant of the MCHRI gene or a susceptibility to the disorder. The MCHRI protein or polymnucleotide is useful for preparing a medicament for treating or preventing obesity related to the presence of a molecular variant of the MCHRI gene. This polymnucleotide represents the melanin-concentrating hormone receptor 1 locus clone of the invention. This sequence is not shown in the specification. It has been taken from the Genbank accession number Z86090 provided in the specification.
 Sequence 122557 BP; 34723 A; 28259 C; 27523 G; 32052 T; 0 U; 0 Other;
 Query Match 37.1%; Score 148.4; DB 12; Length 122557;
 Best Local Similarity 67.8%; Pred. No. 1.2e-30;
 Matches 221; Conservative 1; Mismatches 102; Indels 2; Gaps 1;
 24 GCATGGAACCCAAATTAATAAGATTTGACGCCAGCAGACATCGCTGAATG 83
 67983 GCTGCAGAGGCAAGAGATTGTAATAAGGCCAGGCCGCGTGCACCA 68042
 84 CCGTAATCCAGCACTTCGGGAGGCCCAAGGTGGGATCACTGAGTCAAGATCG 143
 68043 CCGTAATCCAGCACTTCGGGAGGCCCAAGGTGGGATCACTGAGTCAAGATCG 68102
 144 AGACCATCTGCGCAACATGTGTAAACCCGCTTTACTTAATAAATACAAAAATAGCTGG 203
 68103 AGACCATCTGCGCAACATGTGTAAACCCGCTTTACTTAATAAATAGCTGG 68162
 204 GCATGTGGGACACACCTGTAGTCCCACTACTCAGAGCGCGGAGATTCAGTGAAGCTGA 263
 68163 GCATGTGGGACACACCTGTAGTCCCACTACTCAGAGCGCGGAGATTCAGTGAAGCTGA 68222
 264 GATGCAGAGTAGAGCCGAATATCAGATCAGAGA--GTGACAGAGTAGAGCKCGTCTC 321
 68223 GAACCCGGAGGACAGAGTTGCAATGGGCGCAGATGTGCAAGATGAAATCTCCCTCTC 68282
 322 AAAAAACAACAACAAAAACAAAAA 347
 68283 AAAAAAAGAAAGAAAGAAA 68308
 RESULT 50
 AAK79514
 ID AAK79514 standard; DNA; 9469 BP.
 AC AAK79514;
 XX

DT 07-NOV-2001 (first entry)
XX Human immune/haematopoietic antigen genomic sequence SEQ ID NO:34326.
DE
XX Human; immune; haematopoietic; immune/haematopoietic antigen; cancer;
KM cytostatic; gene therapy; vaccine; metastasis; ds.
XX
OS Homo sapiens.
XX
PN MO200157182-A2.
XX
PD 09-AUG-2001.
XX
PF 17-JAN-2001; 2001MO-US001354.
XX
XX 31-JAN-2000; 2000US-0179065P.
PR 04-FEB-2000; 2000US-0180628P.
PR 24-FEB-2000; 2000US-0184664P.
PR 02-MAR-2000; 2000US-0186350P.
PR 16-MAR-2000; 2000US-0189874P.
PR 17-MAR-2000; 2000US-0190076P.
PR 18-APR-2000; 2000US-0198123P.
PR 19-MAY-2000; 2000US-0205515P.
PR 07-JUN-2000; 2000US-0209467P.
PR 28-JUN-2000; 2000US-0214886P.
PR 30-JUN-2000; 2000US-0215135P.
PR 07-JUL-2000; 2000US-0216647P.
PR 07-JUL-2000; 2000US-0216880P.
PR 11-JUL-2000; 2000US-0217487P.
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PR 14-JUL-2000; 2000US-0218290P.
PR 26-JUL-2000; 2000US-0220963P.
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PR 14-AUG-2000; 2000US-0224519P.
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PR 14-AUG-2000; 2000US-0225266P.
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PR 14-AUG-2000; 2000US-0225757P.
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PR 18-AUG-2000; 2000US-0226279P.
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PR 01-SEP-2000; 2000US-0229287P.
PR 01-SEP-2000; 2000US-0229343P.
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PR 08-NOV-2000; 2000US-0246474P.
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PR 05-DEC-2000; 2000US-0251030P.
PR 05-DEC-2000; 2000US-0251988P.
PR 05-DEC-2000; 2000US-0256719P.
PR 06-DEC-2000; 2000US-0251479P.
PR 08-DEC-2000; 2000US-0251856P.
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PR 11-DEC-2000; 2000US-0254097P.
PR 05-JAN-2001; 2001US-02559678P.

XX (HUMA-) HUMAN GENOME SCI INC.

XX Rosen CA, Barash SC, Ruben SM;

XX WPI; 2001-483426/52.

XX Nucleic acids encoding human immune/hematopoietic antigen polypeptides.
XX useful for preventing, diagnosing and/or treating cancers and metastasis.

XX Disclosure: SEQ ID NO 34326; 3071pp + Sequence Listing; English.

XX AAK54951 to AAK64702 encode the human immune/hematopoietic antigen (I)
CC amino acid sequences given in AAM82170 to AAM91921. (I) have cytosolic
CC activity, and can be used in gene therapy and vaccine production. (I)
CC proteins and polynucleotides may be used in the prevention, diagnosis and
CC treatment of diseases associated with inappropriate (I) expression. For
CC example, they may be used to treat disorders associated with decreased
CC expression by rectifying mutations or deletions in a patient's genome
CC that affect the activity of (I) by expressing inactive proteins or to
CC supplement the patient's own production of (I). Additionally, (I) may be
CC polynucleotides may be used to produce the secreted (I), by inserting the
CC nucleic acids into a host cell and culturing the cell to express the
CC protein. (I) proteins and polynucleotides may be used to prevent,
CC diagnose and treat immune/hematopoietic-related diseases, especially
CC cancers and cancer metastases of hematopoietic-derived cells. AAK64703
CC to AAK67694 represent human immune/hematopoietic antigen genomic
CC sequences from the present invention. AAK54942 to AAK54950 and AAM82169
CC represent sequences used in the exemplification of the present invention
XX

XX Sequence 9469 BP; 2940 A; 2203 C; 1941 G; 2385 T; 0 U; 0 Other;

XX Query March 37.0%; Score 148; DB 4; Length 9469;

XX Best Local Similarity 74.2%; Pred. No. 7.1e-31;

XX Matches 187; Conservative 0; Mismatches 65; Indels 0; Gaps 0;

QY 46 TAAACATTGTCAGGCGAGCATGACACTGGCTGTAATGCTTAATCCAGCACTTCGGG 105

DB 1141 TATGACGAAATAGTGTGCGGCGAGGTGCTCATGCTTAATCCAGCACTTTGGG 1200

QY 106 AGGCCAAGTGGGCGGATCACTTGAAGTCAAGAGATCGAACATCTGGCCAACTGCT 165

DB 1201 AGGCCAAGTGGGAGATCACTTGAAGTCAAGAGATCGAACATCTGGCCAACTGCT 1260

QY 166 GAAACCCCGTCTTAACTAAATAAATAAATAAATAAATAAATAAATAAATAAATAA 225

DB 1261 GAAACCCCGTCTTAACTAAATAAATAAATAAATAAATAAATAAATAAATAAATAA 1320

QY 226 TCCAGAGTACTCAGGAGCGGAGATTGCAAGTGAAGTGAAGTGAAGCGGAATC 285

DB 1321 TCCAGAGTACTCAGGAGCGGAGATTGCAAGTGAAGTGAAGTGAAGCGGAATC 1380

QY 286 ACAAGTCAAGA 297

DB 1381 AGTGAGCAAGA 1392

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Job time : 542 secs

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GenCore version 5.1.9
Copyright (c) 1993 - 2006 Bioceleration Ltd.

OW nucleic - nucleic search, using sw model

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Title:

SEQ1-47502C

Perfect score:

399.6

Sequence: 1 ccaggtactcagcactgtgc.....tatgcagagaccacaaaag 400

Scoring table:

IDENTITY_NUC
Gapop 10.0 , Gapext 1.0

Searched: 48236798 seqs, 2795965780 residues

Total number of hits satisfying chosen parameters: 96473596

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%
Listing first 150 summaries

Database :

EST:
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2: gb_est3:*
3: gb_est4:*
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5: gb_est6:*
6: gb_hic:*
7: gb_est2:*
8: gb_est7:*
9: gb_est8:*
10: gb_est9:*
11: gb_g881:*
12: gb_g882:*
13: gb_g883:*
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Pred. No. is the number of results predicted by chance to have a
score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

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1	153	38.3	598	11	AQ416484
2	151.4	37.9	457	11	AQ936397
3	151.4	37.9	617	5	CK330465
4	151.2	37.8	801	3	BU568469
5	151	37.8	357	1	AA405549
6	151	37.8	401	7	BE149224
7	149.6	37.4	513	8	CN272188
8	148.6	37.4	613	13	CZ455807
9	148.6	37.2	542	13	DB137291
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22	147	36.8	589	9	DA808834
23	147	36.8	1085	2	BMS42252
24	146.8	36.7	536	11	AQ381551
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26	146.6	36.7	469	1	AI753536
27	146.6	36.7	490	1	AI434037
28	146.6	36.7	667	14	AG156377
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98	144.4	36.1	1052	3	BM560673	BM560673 AGENCOURT
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101	144.2	36.1	692	14	AG146558	AG146558 Pan trogl
102	144.2	36.1	938	3	BU527958	BU527958 AGENCOURT
103	144	36.0	373	3	AA654778	AA654778 nt73901.s
104	144	36.0	716	13	C2455149	C2455149 MCF737J03
105	144	36.0	895	2	BM452899	BM452899 AGENCOURT
106	143.8	36.0	647	14	AG048965	AG048965 Pan trogl
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108	143.6	35.9	324	10	DM412367	DM412367 HHAGE0130
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110	143.6	35.9	612	1	AA133332	AA133332 zn29202.s
111	143.6	35.9	644	3	BO183419	BO183419 UT-H-EU0-
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114	143.6	35.9	694	11	AQ351427	AQ351427 RPCI11-11
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140	142.8	35.7	610	4	CA442144	CA442144 UI-H-D10-
141	142.6	35.7	1945	6	BC035612	BC035612 Homo sapi
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ALIGNMENTS

RESULT 1
LOCUS AQA16484 598 bp DNA linear GSS 23-MAR-1999
DEFINITION RPCI-11-153H2.TJ RPCI-11 Homo sapiens genomic clone RPCI-11-153H2,
ACCESSION AQA16484
KEYWORDS AQA16484.1 GI:4470608
SOURCE GSS.
ORGANISM Homo sapiens (human)

93	144.4	36.1	301	3	BU957990	BU957990 AGENCOURT
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95	144.4	36.1	505	9	DB278032	DB278032 DB278032
96	144.4	36.1	735	9	DN994393	DN994393 TC111938
97	144.4	36.1	1004	14	DUT99349	DUT99349 f.v01 fp00
98	144.4	36.1	1052	3	BM560673	BM560673 AGENCOURT
99	144.2	36.1	500	14	AG028900	AG028900 Homo sapi
100	144.2	36.1	552	9	DB360773	DB360773 DB360773
101	144.2	36.1	692	14	AG146558	AG146558 Pan trogl
102	144.2	36.1	938	3	BU527958	BU527958 AGENCOURT
103	144	36.0	373	3	AA654778	AA654778 nt73901.s
104	144	36.0	716	13	C2455149	C2455149 MCF737J03
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106	143.8	36.0	647	14	AG048965	AG048965 Pan trogl
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110	143.6	35.9	612	1	AA133332	AA133332 zn29202.s
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112	143.6	35.9	662	14	AG037584	AG037584 Pan trogl
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115	143.6	35.9	673	14	AG143386	AG143386 Pan trogl
116	143.6	35.9	990	12	B2601193	B2601193 WHADAB8TR
117	143.6	35.9	1035	2	BM543345	BM543345 AGENCOURT
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119	143.4	35.9	485	4	CA419549	CA419549 UT-H-FH0-
120	143.4	35.9	693	14	AG179363	AG179363 Pan trogl
121	143.4	35.9	718	8	CR789371	CR789371 DKFZP459J
122	143.4	35.9	831	2	BT758533	BT758533 603022857
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141	142.6	35.7	1945	6	BC035612	BC035612 Homo sapi
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147	142.2	35.6	575	7	AV733228	AV733228 AV733228
148	142.2	35.6	761	9	DA834202	DA834202 DA834202
149	142	35.5	279	1	AT081241	AT081241 oy67b01.x
150	142	35.5	360	1	AF236698	AF236698 AF236698

REFERENCE	Mammalia: Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Eumalia; Eutheria; Eucarchontoglires; Primates; Catarrhini; Homiinae; Homo.		
AUTHORS	Zhao, S., Adams, W.D., Nierman, W., Malek, J., de Jong, P. and Venter, J. C.		
TITLE	Use of BAC End Sequences from Library RPCI-11 for Sequence-Ready Map Building		
JOURNAL	Unpublished (1997)		
COMMENT	Other_GSSE: RPCI11-153H2_TV Contact: Shaying Zhao, William Nierman, Mark Adams Department of Eukaryotic Genomics The Institute for Genomic Research 9712 Medical Center Dr., Rockville, MD 20850 Tel: 301 838 0200 Fax: 301 838 0208 Email: hbe@tigr.org Clones are derived from the human BAC library RPCI-11. For BAC library availability, please contact Pieter de Jong (pieter@tigr.org, med.buffalo.edu). Clones may be purchased from BACPAC Resources (http://bacpac.med.buffalo.edu/ordering) or from Research Genet cs (info@resgen.com). BAC end search page: http://www.tigr.org/tcdb/humgen/bac_end_search/bac_end_search.html. Seq primer: SP6 Class: BAC ends.		
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DB	150	CCTGTAATCCAGACACTTCGGGAGGCCAGGTGGCGGATGACCTGAGGTGAGAGTTTG	209
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QY	204	GCAATGTGTGGACACACCTGTGTAGTCCCACTACTCAGAGGCCGGAGATGACAGTGA	263
DB	270	GCGTGTGTGTGCACGCGCTATATATCCAGCTACTCGGGAGGCGAGAGTTGCAGTGAGCAGA	329
QY	264	GATCGCAGAGTGAAGCCGAATTCACAGATTCACAGAGTGAAGAGTGAAGACCCGCTTCAA	323
DB	330	GATTG-----TGCATTTGCAAGCTGACCTGGGGCAACAAGATGAACCTGTGTCTAA	381
QY	324	AAACACACACAAAAACAAAAAACCATAGACATTG	360
DB	382	AAATAATATATATATATATATATATACACAGATCAAGTAGT	418
RESULT 2	AG936397 457 bp DNA linear GSS 15-MAR-2001		
LOCUS	RPCI-11-Seg1-4-200N9-T7 RPCI-11 Human segment 1-4 genomic library		
DEFINITION	Homo sapiens genomic clone RPCI-11-Seg1-4-200N9-T7 similar to Human Chr 14q24.3, genomic survey sequence.		
ACCESSION	AG936397		
VERSION	AG936397.1 GI:6654471		
KEYWORDS	GSS.		

SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini; Homnidae; Homo.
REFERENCE 1 (bases 1 to 457)
 Osoegawa, K., Mammoser, A.G., Wu, C., Frengen, E., Zeng, C., Caranese, J.J. and de Jong, P.J.
 A bacterial artificial chromosome library for sequencing the complete human genome
 Genome Res. 11 (3), 483-496 (2001)
JOURNAL 11230172
PUBMED
COMMENT Contact: de Jong, P.J.
 Children's Hospital Oakland Research Institute
 747 Fifty Second Street, Oakland, CA 94609-1809, USA
 Tel: 510 450 7911
 Fax: 510 450 7924
 Email: pdejong@mail.cho.org
 BAC end sequences. For clone availability please contact Pieter de Jong (pdejong@mail.cho.org). BACPAC Resources WWW site: www.choi.org/bacpac
 Seq primer: 17
 Class: BAC ends.

FEATURES
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ORIGIN
 Query Match 37.9%; Score 151.4; DB 11; Length 457;
 Best Local Similarity 75.8%; Pred. No. 4e-19;
 Matches 213; Conservative 1; Mismatches 56; Indels 11; Gaps 2;

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 DB 47 ATGCTGTAAATCCAGCACTTCGGAGGCGCGAGTGGCGAGATCACTGAGGTCAAGA 106
 OY 141 TCGAGACATCTGCGCAATGTTGAACCCCGCTTACTAAATAACAAATAATGC 200
 DB 107 TCGAGACATCTGCGCAATGTTGAACCCCGCTTACTAAATAACAAATAATGC 166
 OY 201 TGGGCATGTGGCACAACCTGTAGTCCAGCTACTCAGAA-----GCCGAGATTGC 253
 DB 167 TGGGCATGTGGTATGATCTGTAGTCCAGCTACTCAGAGGCTGAGGCAAGAGTTG 226
 OY 254 AGTAGAGTAGATCGCAGAGTGAAGCGGAATCAAGAT---CACAGAGTGAAGAGATG 309
 DB 227 CTTGAAGCTGAGAGGCAAGAGGTTGCAAGTGAACCAAGATATCTCAGCTTAGAGCAGACG 286
 OY 310 AGAGCCCTCTCAAAAACAACAACAAAAAACAACCA 350
 DB 287 AGACTCGGTCTCAAAAAAANNAANNAACACCTA 327

RESULT 3
 LOCUS CK430465 617 bp mRNA linear EST 06-JAN-2004
 DEFINITION OJ49G05.Y1 Human lacrimal gland, unamplified: OJ Homo sapiens cDNA clone OJ49G05 5', mRNA sequence.
 ACCESSION CK430465
 VERSION CK430465.1 GI:40678584
 KEYWORDS EST.
 SOURCE Homo sapiens (human)
 ORGANISM Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini; Homnidae; Homo.

REFERENCE 1 (bases 1 to 617)
 Ozyildirim, A.M., Wistow, G.J., Gao, J., Wang, J., Dickinson, D.P., Frierson, H.F., Jr and Laurie, G.W.
 The lacrimal gland transcriptome is an unusually rich source of rare and poorly characterized gene transcripts
 Invest. Ophthalmol. Vis. Sci. 46 (5), 1572-1580 (2005)
JOURNAL 1581553
PUBMED
COMMENT Contact: Wistow G
 Section on Molecular Structure and Function
 National Eye Institute
 6/331, NIH, Bethesda, MD 20892-2740, USA
 Tel: 301 402 3452
 Fax: 301 496 0078
 Email: gwen@helix.nih.gov
 Plate: 49 row: 9 column: 05
 Seq primer: M13RP1 reverse primer (ABI).
 Location/Qualifiers

FEATURES
 source
 1..617
 /organism="Homo sapiens"
 /mol_type="mRNA"
 /db_xref="taxon:9606"
 /clone="OJ49G05"
 /issue_type="lacrimal gland"
 /dev_stage="Adult"
 /lab_host="EMD10B"
 /clone_lib="Human lacrimal gland, unamplified: OJ"
 /note="Organ: Eye; Vector: pCMVSPORT6; RNA was extracted from 2 human lacrimal glands. A directionally cloned cDNA library in the pCMVSPORT6 vector (Life Technologies) was constructed at Bioserve Biotechnology (Laurel MD) essentially following the protocols of the SuperScript Plasmid System full details of which are contained in the manufacturer's instruction manual (http://www.lifetech.com/). First strand synthesis was carried out using a Not I primer-adaptor (5'-pGAGTCTCTGATGCGGAGCGCGCCGCTT15-3'). EST analysis was performed on the unamplified library at the NIH Intramural Sequencing Center (NISC)."

ORIGIN
 Query Match 37.9%; Score 151.4; DB 5; Length 617;
 Best Local Similarity 76.4%; Pred. No. 3.7e-19;
 Matches 201; Conservative 1; Mismatches 52; Indels 9; Gaps 1;

OY 81 ATGCTGTAAATCCAGCACTTCGGAGGCGCAAGTGGCGGATCACTGAGGTCAAGA 140
 DB 89 AGCGCTGTAAATCCAGCACTTCGGAGGCGCAAGTGGCGGATCACTGAGGTCAAGA 148
 OY 141 TCGAGACATCTGCGCAATGTTGAACCCCGCTTACTAAATAACAAATAATGC 200
 DB 149 TCGAGACATCTGCGCAATGTTGAACCCCGCTTACTAAATAACAAATAATGC 208
 OY 201 TGGGCATGTGGCACAACCTGTAGTCCAGCTACTCAGAGCGGAGATGAGTAGAG 260
 DB 209 CAGGCATGTGGCACAACCTGTAGTCCAGCTACTCAGAGCGGAGAGCTGCAATAGC 268
 OY 261 TGAAGTGCAGAGTGAAGCGGAATCAAGATCAAGAGTGAAGAGTGAAGAGCTCT 320
 DB 269 CAAGATCATGCTATGCACTCCAGCTTAGGTGA-----CAGAGTAGAGTCTGTCT 319
 OY 321 CAATAACAACAACAAACAA 343
 DB 320 CAATAACAACAAACAAACAA 342

RESULT 4
 LOCUS BUS68469 801 bp mRNA linear EST 16-SEP-2002
 DEFINITION AGENCOURT_1040512 NIH_MGC_82 Homo sapiens cDNA clone IMAGE:661547 5', mRNA sequence.
 ACCESSION BUS68469
 VERSION BUS68469.1 GI:22918758
 KEYWORDS EST.

SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini;
 Hominae; Homo.
REFERENCE 1 (bases 1 to 801)
AUTHORS NIH-MGC <http://mgc.nci.nih.gov/>.
TITLE National Institutes of Health, Mammalian Gene Collection (MGC)
JOURNAL Unpublished (1999)
COMMENT Contact: Robert Strausberg, Ph.D.
 Email: rsb@nih.gov
 Tissue Procurement: CLONTECH
 CDNA Library Preparation: CLONTECH Laboratories, Inc.
 DNA Sequencing by: Agencourt Bioscience Corporation
 Clone distribution: MGC clone distribution information can be
 found through the I.M.A.G.E. Consortium/BLAST at:
<http://image.llnl.gov>
 Plate: L12CM857 row: m column: 19
 High quality sequence stop: 553.
FEATURES
 source
 1..801
 /organism="Homo sapiens"
 /mol_type="mRNA"
 /db_xref="taxon:9606"
 /clone="IMAGE:6615547"
 /lab_host="DH10B (TI phage-resistant)"
 /note="Organ: testis; Vector: pDNR-LIB (Clontech); Site 1:
 SfiI (ggcgccctgcgc); Site 2: SfiI (ggccattggcc); 5' adaptor
 sequence: 5'-CAGCGCCATTATGGCC-3' and 3' adaptor sequence:
 5'-ATTCTAGAGCGCGAGCGCGACGACATG-dT(30)BN-3' (where B = A,
 C, or G and N = A, C, G, or T). Average insert size
 1.35 kb (range 0.9-4.0 kb). 14/15 colonies contained
 inserts by PCR. This library was enriched for full-length
 clones and was constructed by Clontech Laboratories (Palo
 Alto, CA)."
ORIGIN
 Query Match 37.8%; Score 151.2; DB 3; Length 801;
 Best Local Similarity 74.6%; Pred. No. 3.9e-19;
 Matches 206; Conservative 1; Mismatches 59; Indels 10; Gaps 1;
 Db 84 CCTGTAATCCAGCACTTCGGAGGCCAAGGTGGCGGATCATCCTGAGGTCAAGATCG 143
 |||||
 17 CCGTAATCCAGCACTTCGGAGGCCAAGGTGGCGGATCATCCTGAGGTCAAGATCG 76
 |||||
 144 AGACATCTCTGGCCCAACATGTGTGAACCCCGCTTTACTTAAATAATCAAAATAATGTCTGG 203
 |||||
 77 AGACCAAGCTTGGCCCAACATGTGTGAACCCCGCTTTACTTAAATAATCAAAATAATGTCTGG 136
 |||||
 204 GCATGTTGGACACACACCTGTAGTCCAGCTACTTCAGAGCCGGAATTCAGTGGAGTGA 263
 |||||
 137 GCATGTTGGACATGCTCTGTATATCCAGCTTCCAGAGCCGGAATTCAGTGGAGTGA 196
 |||||
 264 GATCGACAGATGAGCCGAATTCACAGATTCACAGATGACAGAGTGAAGACACCCGCTTCAA 323
 |||||
 197 GCTGGCA-----CCATTGCACTCCAGCTCGGATGAGGAGGAGGAGCTTATCTCAA 246
 |||||
 324 AAACACACACCAAAAAACCAAAATCAATTAAGCATT 359
 |||||
 247 AAAAAATAAAAAAATAAATAATTAATAATTAAT 282
 |||||
RESULT 5
 AA405549/c 357 bp mRNA linear EST 17-MAY-1997
 LOCUS zw39f03.r1 Soares total fetus Nb2HF8.9w Homo sapiens CDNA clone
 DEFINITION IMAGE:772445 5' similar to contains Alu repetitive element;; mRNA
 sequence.
 AA405549 AA405549.1 GI:2063141
ACCESSION
VERSION

REFERENCE	AUTHORS
TITLE	Journal
JOURNAL	Comment
COMMENT	Washington University School of Medicine 444 Forest Park Parkway, Box 8501, St. Louis, MO 63108 Tel: 314 286 1800 Fax: 314 286 1810 Email: east@wustl.edu This clone is available royalty-free through LANT; contact the IMAGE Consortium (info@image.llnl.gov) for further information. Seq primer: -28ml3 rev2 ET from Amersham High quality sequence stop: 322. Location/Qualifiers
FEATURES	source
source	1..357 /organism="Homo sapiens" /mol_type="mRNA" /db_xref="taxon:9606" /clone="IMAGE:772445" /dev_stage="8-9 weeks" /lab_host="DH10B" /note="Vector: pRT73D-PacI; Site_1: Not I; Site_2: Eco RI; 1st strand cDNA was prepared from mRNA obtained from pooled 8-9 week (local) fetus material with a Not I - oligo(dT) primer (5' TGTATCCATCTGAAATGAGGAGCGCGCTTAATTTTTTTTTTTT 3'). Double-stranded cDNA was ligated to Eco RI adaptors (Pharmacia), digested with Not I and cloned into the Not I and Eco RI sites of the modified pRT73 vector. Library went through one round of normalization, and was constructed by Bento Soares and M. Fatima Bonaldo."
ORIGIN	
Query Match	37.8%; Score 151; DB 1; Length 357;
Best Local Similarity	77.0%; Pred. No. 5e-19;
Matches	184; Conservative 0; Mismatches 55; Indels 0; Gaps 0;
32	AACCAATATTAATTAAGACATTGTCAGGCGGAGCATACACTGGCTGAATGCCCTGAT 91
287	AATCCGAGATTAAGAAATTAAGCATTAATTTGGTCGGACCGCTGTCAGCGCTGAT 228
92	CCAGCACTTCGGGAGGCGCAAGATGGGGCGATCACTTAGGTCAAGAGATCGAACATC 151
227	CTCAGCACTTTGGGAGGCGCGAGGAGGTGATCACTTAGGTCAAGAGATCGAACATC 168
152	CTGGCAACATGTGAAAACCCGCTTTTACTTAATAATCAAAAATATAGCTGGGATGTTG 211
167	CTGGCAACATGTGAAAACCCGCTTTTACTTAATAATCAAAAATATAGCTGGGATGTTG 210
212	GCACACACCTGTAGCCGACGCTACTCAGAGACCGGAGATTCAGATGAGCTGATCGCA 270
107	GCATGACCTGTATATCCGGCTACTCGGAGAGCTGAGGTTGCAGTGAAGCAAGATTGCA 49
RESULT 6	
LOCUS	BE149224 401 bp mRNA linear EST 21-JUN-2000
DEFINITION	RC3-H10252-120200-014-C06 H10252 Homo sapiens cDNA, mRNA sequence.
ACCESSION	BE149224
VERSION	BE149224.1 GI:8611948
KEYWORDS	EST
SOURCE	Homo sapiens (human)

ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini; Homiidae; Homo.
1 (bases 1 to 401)

REFERENCE
Dias Neto, E., Garcia Correa, R., Verjowski-Almeida, S., Briones, M.R., Nagai, M.A., da Silva, W. Jr., Zago, M.A., Bordin, S., Costa, F.F., Goldman, G.H., Carvalho, A.F., Matsukuma, A., Bala, G.S., Simpson, D.H., Brunstein, A., de Oliveira, P.S., Bucher, P., Jongeneel, C.V., O'Hare, M.U., Soares, F., Brentani, R.R., Reis, L.F., de Souza, S.J. and Simpson, A.J.
Shotgun sequencing of the human transcriptome with ORF expressed sequence tags
Proc. Natl. Acad. Sci. U.S.A. 97 (7), 3491-3496 (2000)

JOURNAL
10737800

COMMENT
Contact: Simpson A.U.G.
Laboratory of Cancer Genetics
Ludwig Institute for Cancer Research
Rua Prof. Antonio Prudente 109, 4 andar, 01509-010, Sao Paulo-SP, Brazil
Tel: +55-11-2704922
Fax: +55-11-2707001
Email: asimpson@ludwig.org.br
This sequence was derived from the FAPESP/LICR Human Cancer Genome Project. This entry can be seen in the following URL:
(http://www.ludwig.org.br/scripts/gethtml2.pl?cl=RC2-HT0252-120
200-014-c06&t3=2000-02-12&t4=1)
Seq primer: puc 18 forward
High quality sequence start: 16
High quality sequence stop: 401.
Location/Qualifiers
1..401
/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/dev_stage="Adult"
/clone_1lb="HT0252"
/note="Organ: head_neck; Vector: puc18; Site 1: Sma1; Site 2: Sma1; A mini-library was made by cloning products derived from ORESTES PCR (U.S. Letters Patent application No. 196,716 - Ludwig Institute for Cancer Research) profiles into the puc 18 vector. Reverse transcription of tissue mRNA and cDNA amplification were performed under low stringency conditions."

ORIGIN
Query Match 37.8%; Score 151; DB 7; Length 401;
Best Local Similarity 78.3%; Pred. No. 4.9e-19;
Matches 206; Conservative 0; Mismatches 55; Indels 2; Gaps 2;

84 CCTGTAATCCAGCACTTCGGAGGCCAAGTGGCGGATCACTGAGGTCAAGATCG 143
42 CCTGTAATCCAGCACTTCGGAGGCCAAGTGGCGGATCACTGAGGTGAGGTTTG 101
144 AGACCATCTGGCCAAATG-GTGAATCCCGCTCTTACTAAATAACAAAAATAGCTG 202
102 AGACCAAGCTGGCCAAATGTTAAATCCCGCTCTTACTAAATAACAAAAATTAAGCA 161
203 GGCATGTTGGACACACCTGTAGTCCAGCTACTCAGAGGACCGAGATTGACAGTGC 262
162 GGCCTGTGGTGGCTGCTGCTGATCCAGCTACTCAGAGGACCGAGATTGACAGTGC 221
263 AGATGACAGAGTGAAGCCGAAATCAAGATCAAGAGTGAAGAGTGAAGACKCCGCTCA 322
222 AGATGACACTACTGACACTAGCTACCTCCAGCTGGGTGA-CAGACAAAGACACTGTCTCA 280

QY 323 AAAACAACAACAAAAA 345
DB 281 AAAACAACAACAAAAA 303

RESULT 7
CN272188

LOCUS CN272188 513 bp mRNA linear EST 16-MAY-2004
DEFINITION 17000600002901 GRN_PREHEP Homo sapiens CDNA 5', mRNA sequence.
ACCESSION CN272188
VERSION CN272188.1 GI:47288602
KEYWORDS EST.
SOURCE
ORGANISM Homo sapiens (human)
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini; Homiidae; Homo.
1 (bases 1 to 513)

REFERENCE
Brandenberger, R., Wei, H., Zhang, S., Lei, S., Murage, J., Fisk, G.J., Li, Y., Xu, C., Fang, R., Guejler, K., Rao, M.S., Mandalam, R., Lebowicki, J. and Stanton, L.W.
Transcriptome characterization elucidates signaling networks that control human ES cell growth and differentiation
Nat. Biotechnol. 22 (6), 707-716 (2004)
15146197

JOURNAL
15146197

COMMENT
Contact: Brandenberger R
Regenerative Medicine
Genon Corporation
230 Constitution Drive, Menlo Park, CA 94025, USA
Tel: 650 473 8658
Fax: 650 473 7760
Email: rbrandenberger@genon.com
Insert Length: 513 Std Error: 0.00.
Location/Qualifiers
1..513
/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/issue_type="embryonic stem cells, DMSO-treated H9 cell line"
/clone_1lb="GRN_PREHEP"
/note="Oligo dt primed, full-length enriched cDNA library from DMSO-treated h9s cell line H9 (p22) maintained in feeder-free conditions"

ORIGIN
Query Match 37.4%; Score 149.6; DB 8; Length 513;
Best Local Similarity 71.4%; Pred. No. 8.6e-19;
Matches 197; Conservative 0; Mismatches 79; Indels 0; Gaps 0;

81 ATGCTGTAATCCAGCACTTCGGAGGCCAAGTGGCGGATCACTGAGGTCAAGAGA 140
51 ATGCTGTAATCCAGCACTTCGGAGGCCAAGTGGCGGATCACTGAGGTCAAGAGA 110
141 TCGAGACATCTGGCCAAATGATGAAACCCCGCTTACTAAATAACAAAAATAGC 200
111 TTGAACCAAGCTGGCCAAATGATGAAACCCCGCTTACTAAATAACAAAAATAGC 170
201 TGGGATGTTGGACACACCTGTAGTCCAGCTACTCAGAGCGGAGATTGACAGTGC 260
171 TGGGATGTTGGACAGTGCCTGTAGTCCAGCTACTCAGAGCGGAGATTGACAGTGC 230
261 TGAATGTCAGAGTGAAGCCGAAATCAAGATCAAGAGTGAAGAGTGAAGACKCCGCT 320
221 TGAGGTGTGTCACACTGCTCAGCTTGGTGAAGAGTGAAGAGTGTCTCAAAAAA 290
321 CAAAAACAACAACAAAAA 356
291 AAAAAACAACAACAAAAA 326

QY 321 CAAAAACAACAACAAAAA 356
DB 291 AAAAAACAACAACAAAAA 326

RESULT 8
CN2455807
LOCUS CN2455807 613 bp DNA linear GSS 20-OCT-2005
DEFINITION MCF73H04TF Human MCF7 breast Cancer cell line library (MCF7_1)
ACCESSION CN2455807
VERSION CN2455807.1 GI:77932102
KEYWORDS GSS.
SOURCE Homo sapiens (human)

ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini; Homiidae; Homo.

REFERENCE 1 (bases 1 to 613)
Volik S.V., Raphael B.J., Huang G.-Q., Murnane J., Brebner J.H., Bajaratowicz K., Paris P., Tao Q., Kowbel D., Lapuk A.V., Kuo W.-L., Shagin D.A., Shagina I.A., Magrane G., Gray J.W., Jan F.-C., de Jong P., Pezner P. and Collins C.
Decoding the genomic architecture and high throughput detection of fusion transcripts in breast cancer cell lines: implications for a tumor genome project
Unpublished (2005)

JOURNAL
COMMENT
Contact: Volik SV
Colin Collins' lab
UCSF Comprehensive Cancer Center
UCSF Box 0808, San Francisco, CA 94143-0808, USA
Tel: 415 502 7066
Fax: 415 502 5665
Email: svolik@cc.ucsf.edu
This clone is available from Amplicon Express
http://www.genomex.com
Class: BAC ends.

FEATURES
source
1. 613
/organism="Homo sapiens"
/mol_type="genomic DNA"
/db_xref="taxon:9606"
/clone="MCF7_38_H04"
/sex="female"
/clone_lib="Human MCF7 breast cancer cell line library (MCF7_1)"
/note="Vector: pBCBAC1; Site_1: HindIII; This library was constructed from MCF7 breast cancer cell line by Amplicon Express (http://www.genomex.com) using their standard procedure."

ORIGIN

Query Match 37.4%; Score 149.6; DB 13; Length 613;
Best Local Similarity 75.6%; Pred. No. 8.3e-19;
Matches 201; Conservative 1; Mismatches 55; Indels 9; Gaps 1;

81 ATGGCTGTAATCCCGACACTTCGGAGGCGCAAGTGGGGGATGTCAGGTCAAGGA 140
87 ACGCTGTAATCTTGAAGCTTTGGAGGCGCAAGTGGATGCTTGAAGTCAAGGT 146
141 TCGAGACCATCTGGCCAAATGTGAACCCCGTCTTACTAAATAACAAAAATAGC 200
147 TCGAGACCATCTGCTAATCAATGTGAACCCCGTCTTACTAAATAACAAAAATAGT 206
201 TGGGATGTTGGACACACACTGTAGTCCAGTACTCAGAGCCGGAATTGCACTGAGC 260
207 TGGACATCTGGCAGACCGCTGTAGTCCAGTCCCTGGAGTGGAGCTTGCAGTGAAC 266
261 TGAGATGGCAGAGTGAAGCGAAATACAGATCAGAGATGAGAGAGAGACCGGCT 320
267 CGAGATCAGCGCACTGCATCCACCTGGGTGA-----CAGAGTGAAGCTGTCT 317
321 CAAAAACAACAACAAAAACAACAAAA 346
318 CAAAAAAGAAAAAGAAAAAGAAAA 343

RESULT 9
LOCUS DB313291 542 bp mRNA linear EST 04-DEC-2005
DEFINITION DB313291 CTONG2 Homo sapiens cDNA clone CTONG2000516 3', mRNA
ACCESSION DB313291
VERSION DB313291.1 GI:83195301
KEYWORDS EST.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens

REFERENCE 1 (bases 1 to 542)
Kimura K., Wakamatsu A., Suzuki Y., Ota T., Nishikawa T., Yamashita R., Yamamoto J., Sekine M., Tsuritani R., Wakaguri H., Ishii S., Sugiyama T., Saito K., Isono Y., Irie R., Kushiida N., Yoneyama T., Otsuka R., Kanda K., Yokoi T., Kondo H., Wagetsuna M., Murakawa K., Ishida S., Ishibashi T., Takahashi Fujii A., Tanase T., Nagai K., Kikuchi H., Nakai K., Isogai T. and Sugano S.
Diversification of Transcriptional Modulation: Large-scale Identification and Characterization of Putative Alternative Promoters of Human Genes
Genome Res. 16 (1), 55-65 (2006)

JOURNAL
PUBMED
16344560
COMMENT
Contact: Takao Isogai
FLJ Project (HRI Team)
Helix Research Institute
2-6-7 Kazusa-Kamatari, Kisarazu, Chiba, 292-0818, Japan
Tel: 81-438-52-3975
Fax: 81-438-52-3986
Email: flj-cdna@nifty.com
NEBO human cDNA project (New Energy and Industrial Technology Developmental Organization, Japan): cDNA library construction: HRI, Research Association for Biotechnology (RAB) and Biotechnology Center, National Institute of Technology and Evaluation; 3'-end one pass sequencing: RAB.

FEATURES
source
1. 542
/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/clone="CTONG2000516"
/tissue_type="tongue, tumor tissue"
/clone_lib="CTONG2"
/note="Vector: pME18SFLJ3"

ORIGIN

Query Match 37.2%; Score 148.6; DB 9; Length 542;
Best Local Similarity 79.5%; Pred. No. 1.3e-18;
Matches 175; Conservative 0; Mismatches 45; Indels 0; Gaps 0;

83 GCTGTATATCCACACACTTGGGAGGCGCAAGTGGGGGATCACCTGAGGTCAAGATC 142
506 GCTGTATATCCACACACTTGGAGGCGCAAGTGGGATCACCTGAGGTCAAGATC 447
143 GAGACCATCTGGCCAAATGTGAACCCCGTCTTACTAAATAACAAAAATAGCTG 202
446 AAGACACAGCTGGCCAAATGTGAACCCCGTCTTACTAAATAACAAAAATAGCTG 387
203 GGCATGTTGGACACACACTGTAGTCCAGTACTCAGAGCCGGAATTGCACTGAGC 262
386 GGCCTGTGGGACATGCTGTAGTCCAGGCTATTGGAGGCTGAGGCTGCAAGTGAAC 327
263 AGATCGCAGAGTGAAGCGAAATCAGATCAGAGTGAAG 302
326 AGATTGGCGCACTGCATCCACCTGGGCGCACAGATGAAG 287

RESULT 10
LOCUS A0541320 614 bp DNA linear GSS 19-MAY-1999
DEFINITION RPCI-11-343C15.TV RPCI-11 Homo sapiens genomic clone
ACCESSION A0541320
VERSION A0541320.1 GI:4871776
KEYWORDS GSS.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini; Homiidae; Homo.

REFERENCE 1 (bases 1 to 614)
 AUTHORS Zhao, S., Adams, M.D., Niernman, W., Malek, J., de Jong, P. and Venter, J.C.
 TITLE Use of BAC End Sequences from Library RPCI-11 for Sequence-Ready Map Building
 JOURNAL Unpublished (1997)
 COMMENT Other GSSs: RPCI-11-343C15.TJ
 Contact: Shaoying Zhao, William Niernman, Mark Adams
 Department of Eukaryotic Genomics
 The Institute for Genomic Research
 9712 Medical Center Dr., Rockville, MD 20850
 Tel: 301 838 0200
 Fax: 301 838 0208
 Email: hbe@tigr.org

Clones are derived from the human BAC library RPCI-11. For BAC library availability, please contact Pieter de Jong (pieter@edlong.med.buffalo.edu). Clones may be purchased from BACPAC Resources (<http://bacpac.med.buffalo.edu/ordering>) or from Research Genet cs (info@resgen.com). BAC end search page: http://www.tigr.org/tdb/hungen/bac_end_search/bac_end_search.html.
 Seq primer: 17
 Class: BAC ends.

FEATURES

source 1..614
 Location/Qualifiers
 /organism="Homo sapiens"
 /mol_type="genomic DNA"
 /db_xref="GDB:7631390"
 /db_xref="taxon:9606"
 /clone="RPCI-11-343C15"
 /sex="Male"
 /cell_type="Lymphocytes"
 /clone_lib="RPCI-11"
 /note="Vector: pBACe3.6; Site_1: EcoRI; Site_2: EcoRI; RPCI11 Human Male BAC Library"

ORIGIN

Query Match 37.1%; Score 148.4; DB 11; Length 614;
 Best Local Similarity 72.1%; Pred. No. 1.4e-18;
 Matches 209; Conservative 1; Mismatches 72; Indels 8; Gaps 1;
 84 CCTGTAATCCAGCACTTCGGAGGCGCAAGGTGGGGGATCACTGAGGTCAAGAGATCG 143
 447 CCTGTAATCCAGCACTTCGGAGGCGCAAGGTGGGGGATCACTGAGGTCAAGATTTG 388
 144 AGACCATCTGGCCCAACATGTGTAACCCCGCTTTACTTAATAAATAACAAAAATAGCTGG 203
 387 AGACCAAGCTGGCCCAACATGTGTAACCCCGCTTTACTTAATAAATAACAAAAATAGCTGG 328
 204 GCATGATGGACACACCTGTAGTCCAGCTACTCAAGAGCGGAGATTGCACTGAGCTGA 263
 327 GCGTGTGGTGGGTGCTGTGTAATCCAGCTACTCAAGAGCGGAGATTGCTAGTGAAC 271
 264 GATCGCAGAGTGAGCGCAATATCAGATCAAGAGTGAGCAGAGTGAAGACKCCGCTCTCA 323
 270 -----CAAAGATTGACACACCTGCACTCCAGTCTGGGGCAGAGAGAGATCTCATCTCA 216
 324 AAACAACAACAAAAAACCAATTAAGACATTTGCTGCGGTT 373
 215 AAAAAAAAAAGAAAAAGAAAAAGAAACATACCTGAATTTTAGGGGT 166

RESULT 11
 DA572262/c
 LOCUS DA572262 847 bp mRNA linear EST 07-NOV-2005
 DEFINITION DA572262 HEMBI Homo sapiens cDNA clone HEMBI100862.5', mRNA
 sequence.
 ACCESSION DA572262
 VERSION DA572262.1 GI:81139202
 KEYWORDS EST.
 SOURCE Homo sapiens (human)
 ORGANISM Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini;

REFERENCE 1 (bases 1 to 847)
 AUTHORS Kimura, K., Wakamatsu, A., Suzuki, Y., Ota, T., Nishikawa, T., Yamashita, R., Yamamoto, J., Sekine, M., Teuriltani, K., Wakaquri, H., Ishii, S., Sugiyama, T., Saito, K., Isono, Y., Irie, R., Koshida, N., Yoneyama, T., Otsuka, R., Kanda, K., Yokoi, T., Kondo, H., Magatawma, M., Murakawa, K., Ishida, S., Ishibashi, T., Takahashi-Fujii, A., Tanase, T., Nagai, K., Kikuchi, H., Nakai, K., Isogai, T. and Sugano, S.
 TITLE Diversification of Transcriptional Modulation: Large-scale Identification and Characterization of Putative Alternative Promoters of Human Genes
 JOURNAL Genome Res. 16 (1), 55-65 (2006)
 PUMED 16344560
 COMMENT Contact: Takao Isogai
 Fij Project (HRI Team)
 Helix Research Institute
 2-6-7 Kazusa-Kamatari, Kisarazu, Chiba, 292-0818, Japan
 Tel: 81-438-52-3975
 Fax: 81-438-52-3986
 Email: fli-cdn@hri.fy.com

HRI human cDNA project; 5'- & 3'-end one pass sequencing; Helix Research Institute (HRI); cDNA library construction; Department of Virology, Institute of Medical Science, University of Tokyo, and HRI.

FEATURES

source 1..847
 Location/Qualifiers
 /organism="Homo sapiens"
 /mol_type="mRNA"
 /db_xref="taxon:9606"
 /clone="HEMBI100862"
 /issue_type="whole embryo, mainly body"
 /dev stage="embryo, 10 weeks"
 /clone_lib="HEMBI"
 /note="Vector: pME18SFL3"

ORIGIN

Query Match 37.1%; Score 148.4; DB 9; Length 847;
 Best Local Similarity 76.8%; Pred. No. 1.3e-18;
 Matches 195; Conservative 1; Mismatches 52; Indels 6; Gaps 1;
 84 CCTGTAATCCAGCACTTCGGAGGCGCAAGGTGGGGGATCACTGAGGTCAAGAGATCG 143
 530 CCTGTAATCCAGCACTTCGGAGGCGCAAGGTGGGGGATCACTGAGGTCCGAGTTCA 471
 144 AGACCATCTGGCCCAACATGTGTAACCCCGCTTTACTTAATAAATAACAAAAATAGCTGG 203
 470 AGACCAAGCTGGCCCAACATGTGTAACCCCGCTTTACTTAATAAATAACAAAAATAGCTGG 411
 204 GCATGATGGACACACCTGTAGTCCAGCTACTCAAGAGCGGAGATTGCACTGAGCTGA 263
 410 GCATGATGGACACACCTGTAGTCCAGCTACTCAAGAGCGGAGATTGCACTGAGCTGA 351
 264 GATCGCAGAGTGAGCGCAATATCAGATCAAGAGTGAGCAGAGTGAAGACKCCGCTCTCA 323
 350 GATCGCGCAGTGTGCTGCACTCA-----GCTGGGTGACAGAGTGAAGATCTCATCA 297
 324 AAACAACAACAAA 337
 296 AAAAAAAAAAAAA 283

RESULT 12
 AG170656/c
 LOCUS AG170656 691 bp DNA linear GSS 09-JAN-2002
 DEFINITION Pan troglodytes DNA, clone: RP43-039L04.TJ, genomic survey
 sequence.
 ACCESSION AG170656
 VERSION AG170656.1 GI:16700334
 KEYWORDS GSS.
 SOURCE Pan troglodytes (chimpanzee)
 ORGANISM Pan troglodytes
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini;

REFERENCE 1 Hominidae; Pan.
 AUTHORS Fujiyama, A., Hattori, M., Toyoda, A., Taylor, T.D., Yada, T.,
 TITLE Toki, Y., Watanabe, H. and Sakaki, Y.
 JOURNAL BAC end sequences of Library RPCI-43
 COMMENT Unpublished
 REFERENCE 2 (bases 1 to 691)
 AUTHORS Fujiyama, A., Hattori, M., Toyoda, A., Taylor, T.D., Yada, T.,
 TITLE Toki, Y., Watanabe, H. and Sakaki, Y.
 JOURNAL Direct Submission
 Submitted (02-AUG-2001) Asao Fujiyama, The Institute of Physical
 and Chemical Research (RIKEN), Genomic Sciences Center (GSC),
 1-7-22 Suehiro-cho, Tsurumi-ku, Yokohama, Kanagawa 230-0045, Japan
 (E-mail: chimpansecgsc.riken.go.jp, URL: http://nsp.gsc.riken.go.jp/,
 Tel: 81-45-503-9111, Fax: 81-45-503-9170)
 Clones are derived from the chimpanzee BAC library RPCI-43 This BAC
 end was generated during the R&D process and may have higher chance
 of clone tracking errors.
 PRIMERS
 Sequencing: T7
 LIBRARY
 Vector : pBAC3.6
 R.Site 1 : EcoRI
 R.Site 2 : EcoRI.
 Location/Qualifiers
 1. 691
 /organism="Pan troglodytes"
 /mol_type="genomic DNA"
 /db_xref="taxon:9598"
 /clone="RP43-039104.TU"
 /sex="male"
 /cell_type="lymphocytes"
 /clone_id="RPCI-43 Chimpanzee Male BAC Library"
 ORIGIN
 Query Match 37.0%; Score 148; DB 14; Length 691;
 Best Local Similarity 71.1%; Pred. No. 1,7e-18;
 Matches 212; Conservative 1; Mismatches 76; Indels 9; Gaps 1;
 Oy 45 ATAAGACATGTGTCAGGCGGAGGATGACATGCTGTAATGCTGTAATCCAGCACTTGG 104
 Db 405 ATAAAGTTTAAATATGAGCGAGGTGTGTGGTGTGACACCTGTCATCCAGCACTTGG 346
 Oy 105 GAGGCGAAGGTGGGGGATCACCTTGAAGTCAAGAAATGAGACCTCTCTGGCAATGG 164
 Db 345 GAGGTGAGACTGTGGATCGCTTGAAGTCAAGAAATGAGACCTCTGGCAATGG 286
 Oy 165 TGAAGACCCGCTTACTTAAATAATCAAAAAATAGCTGGGCAATGGTGCAACACCTGTA 224
 Db 285 TGAAGCCCTGTCTTACAAAATAATCAAACTTACCTGAGCATGGTGGATGACCTGTA 226
 Oy 225 GTCCAGCTACTCAGAGCCGAGATTCAGTGAAGCTGAGTGCAGAGTGCAGCGGAAT 284
 Db 225 GTCCAGCTACTCAGAGGCTGAGATTCAGTGAAGCTGAGTGCAGATTCAGCTCAG 166
 Oy 285 CACAGATCAGAGTGAAGTGAAGTGAAGTGAAGTGAAGTGAAGTGAAGTGAAGTGAAG 342
 Db 165 CCGTGGCCA-----CAGAGTGAAGTCCGCTCAAAAAAAGAAAAA 117
 RESULT 13
 LOCUS BF965924 749 bp mRNA linear EST 22-JAN-2001
 DEFINITION 602277287P2 NIH_MGC_86 Homo sapiens cDNA IMAGE:4365117 5',
 mRNA sequence.
 ACCESSION BF965924
 VERSION BF965924.1 GI:12333139
 KEYWORDS EST.
 SOURCE Homo sapiens (human)
 ORGANISM Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini;
 Hominidae; Homo.

REFERENCE 1 (bases 1 to 749)
 AUTHORS NIH-MGC http://mgi.nci.nih.gov/
 TITLE National Institutes of Health, Mammalian Gene Collection (MGC)
 JOURNAL Unpublished (1999)
 COMMENT Contact: Robert Strausberg, Ph.D.
 Email: cgabbs-remail.nih.gov
 Tissue Procurement: ATCC
 CDNA Library Preparation: Life Technologies, Inc.
 CDNA Library Arrayed by: The I.M.A.G.E. Consortium (LMNL)
 DNA Sequencing by: Incyte Genomics, Inc.
 Clone distribution: MGC clone distribution information can be
 found through the I.M.A.G.E. Consortium/LMNL at:
 http://image.llnl.gov
 Plate: LMNL0013 row: m column: 22
 High quality sequence stop: 583.
 Location/Qualifiers
 1. 749
 /organism="Homo sapiens"
 /mol_type="mRNA"
 /db_xref="taxon:9606"
 /clone="IMAGE:4365117"
 /tissue_type="osteosarcoma, cell line"
 /lab_host="DH10B (phage-resistant)"
 /clone_id="NIH_MGC_86"
 /note="Organ: bone; Vector: pCMV-SPORT6; Site 1: NotI;
 Site 2: SalI; Cloned unidirectionally; oligo-dT primed.
 Average insert size 1.533 kb. Library enriched for
 full-length clones and constructed by Life Technologies.
 Note: this is a NIH_MGC library."
 ORIGIN
 Query Match 37.0%; Score 148; DB 2; Length 749;
 Best Local Similarity 74.3%; Pred. No. 1.6e-18;
 Matches 199; Conservative 1; Mismatches 66; Indels 2; Gaps 1;
 Oy 81 ATGCTGTAATCCAGCACTTGGGAGGCGCAAGTGGGCGGATCAGCTGAGTCAAGAGA 140
 Db 137 AGCGCTGTAATCCAGCACTTGGGAGGCGTGAAGTGGGAGATCAGCTGAGTGAAGT 196
 Oy 141 TGAAGACCATCTGAGCAACATGATGAAGACCCGCTTTTAAATAATCAAAAAATGAC 200
 Db 197 TAGAGACCATCTGAGCAACATGATGAAGACCCGCTTTTAAATAATCAAAAAATGAC 256
 Oy 201 TGGGATGATGAGCACACCTGTATGCTCCACTCTCAAGAGCCGAGATTCAGTGAAGC 260
 Db 257 CAGCGTGTGTGGCCACATCTGTAGTCCCACTTCTGGGAG--GCTGAAGCAGAGAT 314
 Oy 261 TGAAGTGCAGAGTGAAGCCGAATTCACAGATCAGAGTGAAGTGAAGTGAAGTGAAGT 320
 Db 315 TGCTTGAACCCAGAGGAGGAGGTTCAGTGAAGCTGGGCGACAGAGCAAGACTGTCT 374
 Oy 321 CAAAAACAACAACAAAAAACAAC 348
 Db 375 CAAAAAACAACAAAAAACAAGAAC 402
 RESULT 14
 LOCUS AQ341973 538 bp DNA linear GSS 06-MAY-1999
 DEFINITION RPCI11-111D18.TV RPCI-11 Homo sapiens genomic clone RPCI-11-111D18,
 genomic survey sequence.
 ACCESSION AQ341973
 VERSION AQ341973.1 GI:4166869
 KEYWORDS GSS.
 SOURCE Homo sapiens (human)
 ORGANISM Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini;
 Hominidae; Homo.
 REFERENCE 1 (bases 1 to 538)
 AUTHORS Zhao, S., Adams, M.D., Niernan, W., Malek, J., de Jong, P. and
 Venter, J.C.
 TITLE Use of BAC End Sequences from Library RPCI-11 for Sequence-Ready

JOURNAL
COMMENT
Map Building
Unpublished (1997)
Contact: Shaying Zhao, William Niernman, Mark Adams
Department of Eukaryotic Genomics
The Institute for Genomic Research
9712 Medical Center Dr., Rockville, MD 20850
Tel: 301 838 0200
Fax: 301 838 0208
Email: hbe@tigr.org

Clones are derived from the human BAC library RPC1-11. For BAC library availability, please contact Pieter de Jong (pieter@edlong.med.buffalo.edu). Clones may be purchased from BACAC Resources (http://bacpac.med.buffalo.edu/ordering) or from Research Genetics (info@resgen.com). BAC end search page: http://www.tigr.org/cdb/humgen/bac_end_search.html
Seq primer: 17
Classes: BAC ends.

FEATURES

Location/Qualifiers
1..538
/organism="Homo sapiens"
/mol_type="genomic DNA"
/db_xref="GDB:7542329"
/db_xref="taxon:9606"
/clone="RPC1-11-11D18"
/sex="Male"
/cell_type="Lymphocytes"
/clone_lib="RPC1-11"
/note="Vector: pBACe3.6; Site_1: EcoRI; Site_2: EcoRI; RPC11 Human Male BAC Library"

ORIGIN

Query Match 36.9%; Score 147.6; DB 11; Length 538;
Best Local Similarity 74.6%; Pred. No. 2.1e-18;
Matches 197; Conservative 1; Mismatches 65; Indels 1; Gaps 1;
84 CCTGTAATCCAGCACTTCGAGAGCCAAAGTGGCGGATCACTGAGGTCAAGATCG 143
163 CCTGTAATCCAGCACTTCGAGAGCCAAAGTGGCGGATCACTGAGGTCAAGATCG 222
144 AGACCATCTGGCCAACTGTGTAACCCCGCTCTTAAATAACAAAATATAGCTGG 203
223 AGACCATCTGGCCAACTGTGTAACCCCGCTCTTAAATAACAAAATATAGCTGG 282
204 GCATGTGGCAACCTGTAGTCCAGCTACTCAGAGCCGGAATGCGATGAGCTGA 263
283 GCATGTGGCAACCTGTAGTCCAGCTACTCAGAGCCGGAATGCGATGAGCTGA 341
264 GATCGCAGTGTAGCCGAATACAGATCAAGATGAGAGTGAAGACCCGCTCA 323
342 TGAGCTGAGTGTGCGCACTGCACTTCCAGCTTGGGCAACAGATGAGACTGTCTCA 401
324 AAACCAACAACAAAAA 347
402 AATAATATATATATAGTAATA 425

RESULT 15
AA648957/c 426 bp mRNA linear EST 29-OCT-1997
LOCUS
DEFINITION
n830905.s1 NCI CGAP GCBI Homo sapiens cDNA clone IMAGE:1185176 3' similar to contains Alu repetitive element; contains element PTR5 repetitive element 1; mRNA sequence.
ACCESSION
VERSION
KEYWORDS
SOURCE
ORGANISM
REFERENCE
AUTHORS
TITLE
AA648957
AA648957.1 GI:2575386
EST.
Homo sapiens (human)
Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini; Homnidae; Homo.
NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
National Cancer Institute, Cancer Genome Anatomy Project (CGAP),

JOURNAL
COMMENT
Tumor Gene Index
Unpublished (1997)
Contact: Robert Strusberg, Ph.D.
Email: cgapds-remail.nih.gov
unknown library type
Seq primer: -40m13 fwd. ET from Amersham
High quality sequence stop: 368.

FEATURES

Location/Qualifiers
1..426
/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/clone="IMAGE:1185176"
/tissue_type="germinal center B cell"
/lab_host="DH10B"
/clone_lib="NCI-CGAP GCBI"
/note="Vector: pT7T3D-Pac1; Site_1: Not I; Site_2: Eco RI; 1st strand cDNA was prepared from human tonsillar cells enriched for germinal center B cells by flow sorting (CD20+, IgD-) provided by Dr. Louis M. Staudt (NCI), Dr. David Altman (NCI) and Dr. Gerald Marti (CBER). cDNA synthesis was primed with a Not I - oligo(dT) primer [5'-GTTACCAATCTGAAGGAGCGCGCCCTCATTTTCTTTTCTT-3' (Pharmacia), digested with Not I and cloned into the Not I and Eco RI sites of the modified pT7T3 vector. Library went through one round of normalization, and was constructed by Bento Soares and M. Fatima Bonaldo."

ORIGIN

Query Match 36.9%; Score 147.4; DB 1; Length 426;
Best Local Similarity 71.1%; Pred. No. 2.4e-18;
Matches 224; Conservative 1; Mismatches 82; Indels 8; Gaps 2;
37 AATATATATAGCACTTGTGAGCCAGCACTGACATGACCTGCTGATGCTGATCCAG 96
399 ACATTACAGAAAGAAAGAGTGGCTGCGAGTGTGGCTATGCTGTAATCCAG 340
97 CACTTGGGAGCCCAAGTGGCGGATCACTGAGTCAAGATCGAGACCATCTGGC 156
339 CACTTGGGAGCCCAAGTGGCGGAGTGGCTGAGTCAAGTGGAGACCATCTGGC 280
157 CAACATGTGTAACCCCGCTCTTAAATAACAAAATATAGTGGCATGTGGACA 216
279 CAACATGTGTAACCCCGCTCTTAAATAACAAAATATAGTGGCTGTGGTGTGCA 220
217 CACTGTAGTCCAGCTACTCAGAGCCGAGATTTGAG-----TGAGCTGAGATGCG 269
219 TGCGTGTATCCCGACTTCTGGGAGGCTGAGTGAAGATCACTTAACCCAGAAAGC 160
270 AGAGTGAAGCCGAATACAGATCAAGATGAG-CAGAGTGAAGACCCGCTCAAAAACA 328
159 GGAGGTGCACTGAGCCGAGATCAACCTGGAGAGACAGATGAGACTCCATCTCAAAAAA 100
329 ACAACAAAACAAA 343
99 AAAAAAAAAAAAAA 85

RESULT 16
DR759556 897 bp mRNA linear EST 21-JUL-2005
LOCUS
DEFINITION
HSC4_105.B07.g1 A037 NIH_MGC_262 Homo sapiens cDNA clone IMAGE:7968807 5', mRNA sequence.
ACCESSION
VERSION
KEYWORDS
SOURCE
ORGANISM
REFERENCE
AUTHORS
TITLE
DR759556
DR759556.1 GI:71052256
EST.
Homo sapiens (human)
Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini; Homnidae; Homo.
NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
National Cancer Institute, Cancer Genome Anatomy Project (CGAP),


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RESULT 18
LOCUS      DA180817/c      540 bp      mRNA      linear      EST 01-NOV-2005
DEFINITION DA180817 BRAVY2 Homo sapiens cDNA clone BRAVY2045648 5', mRNA
ACCESSION  DA180817
VERSION     DA180817.1  GI:78555451
KEYWORDS   EST.
SOURCE     Homo sapiens (human)
ORGANISM   Homo sapiens
            Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
            Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini;
            Homnidae; Homo
REFERENCE  1 (bases 1 to 540)
AUTHORS   Kimura,K., Wakamatsu,A., Suzuki,Y., Ota,T., Nishikawa,T.,
            Yamashita,R., Yamamoto,J., Sekine,M., Teuricani,K., Wakaguri,H.,
            Ishii,S., Sugiyama,T., Saito,K., Isono,Y., Irie,R., Kushiida,N.,
            Yoneyama,T., Otsuka,R., Kanda,K., Yokoi,T., Kondo,H., Magatsuma,M.,
            Murakawa,K., Ishida,S., Ishibashi,T., Takahashi-Fujii,A.,
            Tanabe,T., Nagai,K., Kikuchi,H., Nakai,K., Isogai,T. and Sugano,S.
TITLE      Diversification of Transcriptional Modulation: Large-scale
            Identification and Characterization of Putative Alternative
            Promoters of Human Genes
JOURNAL    Genome Res. 16 (1), 55-65 (2006)
PUBMED     16344560
COMMENT    Contact: Takao Isogai
            FLY Project (HRI Team)
            Helix Research Institute
            2-6-7 Kazusa-Kamatari, Kisarazu, Chiba, 292-0818, Japan
            Tel: 81-438-52-3975
            Fax: 81-438-52-3986
            Email: fly-cdna@hri.fly.com
            NEDO human cDNA project (New Energy and Industrial Technology
            Developmental Organization, Japan); cDNA library construction:
            Helix Research Institute (HRI); 5'-end one pass sequencing: HRI,
            Research Association for Biotechnology (Rab) and Biotechnology
            Center, National Institute of Technology and Evaluation; 3'-end one
            pass sequencing: Rab.
FEATURES   Location/Qualifiers
            source          1..540
                        /organism="Homo sapiens"
                        /mol_type="mRNA"
                        /db_xref="taxon:9606"
                        /clone="BRVY2045648"
                        /tissue_type="amygdala"
                        /clone_id="BRVY2"
                        /note="Vector: pME18SFL3"
ORIGIN
Query Match      36.8%; Score 147; DB 9; Length 540;
Best Local Similarity 71.9%; Pred. No. 2.7e-18;
Matches 192; Conservative 0; Mismatches 75; Indels 0; Gaps 0;

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RESULT 19
LOCUS      BG258140/c      545 bp      mRNA      linear      EST 13-FEB-2001
DEFINITION BG258140 NIH_MGC_92 Homo sapiens cDNA clone IMAGE:4510256 5',
            mRNA sequence.
ACCESSION  BG258140
VERSION     BG258140.1  GI:12767956
KEYWORDS   EST.
SOURCE     Homo sapiens (human)
ORGANISM   Homo sapiens
            Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
            Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini;
            Homnidae; Homo.
REFERENCE  1 (bases 1 to 545)
AUTHORS   NIH-MGC http://mgi.nci.nih.gov/.
TITLE      National Institutes of Health, Mammalian Gene Collection (MGC)
JOURNAL    Unpublished (1999)
COMMENT    Contact: Robert Strausberg, Ph.D.
            Email: cgapbs-remail.nih.gov
            Tissue Procurement: ATCC
            cDNA Library Preparation: Life Technologies, Inc.
            DNA Sequencing by: Incyte Genomics, Inc.
            Clone distribution: MGC clone distribution information can be
            found through the I.M.A.G.E. Consortium/LLNL at:
            http://image.llnl.gov
            Plate: LLML0391 row: m column: 09
            High quality sequence stop: 542.
FEATURES   Location/Qualifiers
            source          1..545
                        /organism="Homo sapiens"
                        /mol_type="mRNA"
                        /db_xref="taxon:9606"
                        /clone="IMAGE:4510256"
                        /tissue_type="embryonal carcinoma, cell line"
                        /lab_host="DH10B (phage-resistant)"
                        /note="Organ: testis; Vector: pCMV-SPORT6; Site 1: NotI;
                        Site 2: SalI; Cloned unidirectionally; oligo-dT primed.
                        Average insert size 2.5 kb. Library enriched for
                        full-length clones and constructed by Life Technologies.
                        Note: this is a NIH_MGC Library."
ORIGIN
Query Match      36.8%; Score 147; DB 2; Length 545;
Best Local Similarity 71.9%; Pred. No. 2.7e-18;
Matches 192; Conservative 0; Mismatches 75; Indels 0; Gaps 0;

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LOCUS	DA158439	547 bp	mRNA	linear	EST 30-OCT-2005
DEFINITION	DA158439 BRAMY2 Homo sapiens cDNA clone BAMY2017998 5', mRNA sequence.				
ACCESSION	DA158439				
VERSION	DA158439.1	GI:78296613			
KEYWORDS	EST.				
SOURCE	Homo sapiens (human)				
ORGANISM	Homo sapiens				
	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini; Hominidae; Homo.				
REFERENCE	1 (bases 1 to 547)				
AUTHORS	Kimura,K., Makamatsu,A., Suzuki,Y., Ota,T., Nishikawa,T., Yamashita,R., Yamamoto,J., Sekine,M., Tsutitani,K., Wakaguri,H., Ishii,S., Sugiyama,T., Saito,K., Isono,Y., Irie,R., Kushida,N., Yoneyama,T., Otsuka,R., Kanda,K., Yokoi,T., Kondo,H., Wagaetsuna,M., Murakawa,K., Ishida,S., Isibeshi,T., Takahashi-Fujii,A., Tanase,T., Nagai,K., Kikuchi,H., Nakai,K., Isogai,T. and Sugano,S.				
TITLE	Diversification of Transcriptional Modulation: Large-scale Identification and Characterization of Putative Alternative Promoters of Human Genes				
JOURNAL	Genome Res. 16 (1), 55-65 (2006)				
PUBMED	16344560				
COMMENT	Contact: Takao Isogai FLJ Project (HRI Team) Helix Research Institute 2-6-7 Kazusa-Kamatari, Kisarazu, Chiba, 292-0818, Japan Tel.: 81-438-52-3975 Fax: 81-438-52-3986 Email: flj-cdna@nifty.com NEBO human cDNA project (New Energy and Industrial Technology Developmental Organization, Japan); cDNA library construction: Helix Research Institute (HRI); 5'-end one pass sequencing: HRI, Research Association for Biotechnology (RAB) and Biotechnology Center, National Institute of Technology and Evaluation; 3'-end one pass sequencing: RAB. Location/Qualifiers 1..547 /organism="Homo sapiens" /mol_type="mRNA" /db_xref="taxon:9606" /clone="BRAMY2017998" /tissue_type="amygdala" /clone_id="BRAMY2" /note="Vector: pME18SFL3"				
FEATURES	source				
	ORIGIN				
	Query Match 36.8%; Score 147; DB 9; Length 547; Best Local Similarity 71.9%; Pred. No. 2.7e-18; Matches 192; Conservative 0; Mismatches 75; Indels 0; Gaps 0;				
OY	81 ATGGCTGTAATCCCGCAGCACTTCGGAGGCCAAGGTGGGCGGATCACCTGAGGTCAAAGA 140				
DB	372 ACGCTTAAATCCAGCACTTTGGAGGCTGAAGCCGGTGATCATCTGAAGGTCAAGGAT 313				
OY	141 TCGAGACCATTCTGGCCAAACATGGTGAATCCCGCTTTACTATAAAAAATACAAAATAATAGC 200				
DB	312 TCAAGACCAAGCTGGCCAAACAGTGAATACCCCACTCTTACTATAAAAAATACAAAATAATAGC 253				
OY	201 TGGGCATGTGGCACAACCTGTAGTCCAGTACTACAGAGACCGGAATTGCACTGAGC 260				
DB	252 TGGGATATGTGTGATCATCTGTATATCCAGCTACTTATGGGAGGCTGAAGTTGCACTGAGC 193				
OY	261 TGAGATGGCAAGTGAGCCGAATACAGATACAGATACAGAGTAGAGAGAGCAKCCGCT 320				
DB	192 CAAAGTCCACAATGCACTCCAGCTGGGTGACAGACGAAGATCCGCTCTCCAAAAA 133				
OY	321 CAAAAACAACAACAAAAA 347				
DB	132 TAAATTAATTAATTAATTAATTAATCA 106				
RESULT	21				

DB176347/c	DB176347	569 bp	mRNA	Linear	EST 10-DEC-2005
LOCUS	DB176347	TKIDN2 Homo sapiens cDNA clone	TKIDN2012877 5', mRNA		
DEFINITION	sequence.				
ACCESSION	DB176347				
VERSION	DB176347.1	GI:83514420			
KEYWORDS	EST.				
SOURCE	Homo sapiens (human)				
ORGANISM	Homo sapiens				
REFERENCE	Eukaryote; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini; Homnidae; Homo.				
AUTHORS	1 (bases 1 to 569)				
	Kimura,K., Wakamatsu,A., Suzuki,Y., Ota,T., Nishikawa,T., Yamashita,R., Yamamoto,J., Sekine,M., Tsutitani,K., Makaguri,H., Ishii,S., Sugiyama,T., Saito,K., Isono,Y., Irie,R., Kushida,N., Yoneyama,T., Otsuka,R., Kanda,K., Yokoi,T., Kondo,H., Wagatsuna,M., Murakawa,K., Ishida,S., Ishibashi,T., Takahashi-Fujii,A., Tanase,T., Nagai,K., Kikuchi,H., Nakai,K., Isogai,T. and Sugano,S.				
TITLE	Diversification and Transcriptional Modulation: Large-scale Identification and Characterization of Putative Alternative Promoters of Human Genes				
JOURNAL	Genome Res. 16 (1), 55-65 (2006)				
PUBMED	16344560				
COMMENT	Contact: Takao Isogai FLJ Project (HRI Team) Helix Research Institute 2-6-7 Kanusa-Kamatari, Kisarazu, Chiba, 292-0818, Japan Tel.: 81-438-52-3975 Fax: 81-438-52-3986 Email: flj-cdna@hriity.com				
FEATURES	NEO human cDNA project (New Energy and Industrial Technology Developmental Organization, Japan); cDNA library construction: Helix Research Institute (HRI); 5'-end one pass sequencing: HRI, Research Association for Biotechnology (RAB) and Biotechnology Center, National Institute of Technology and Evaluation; 3'-end one pass sequencing: RAB.				
source	Location/Qualifiers				
	1..569				
	/organism="Homo sapiens"				
	/mol_type="mRNA"				
	/db_xref="taxon:9606"				
	/clone="TKIDN2012877"				
	/tissue_type="Kidney, tumor tissue"				
	/clone_id="TKIDN2"				
	/note="Vector: pME18SFL3"				
ORIGIN					
Query Match	36.8%	Score 147;	DB 9;	Length 569;	
Best Local Similarity	71.9%	Pred. No. 2.7e-16;			
Matches 192;	Conservative 0;	Mismatches 75;	Indels 0;	Gaps 0;	
81	ATGCGTGAATCCGAGCACTTCGGAGGCGCAGGTGGCGGATGACCTGAGTCAAGAGA	140			
386	ACGCTTAATACCCAGACACTTTGGGAGGCTGAGGCCGGTGGATCACTGAGGTCAAGAGT	327			
141	TCGAGACCATCTGCGCCAACTGGTGAATCCCGCTCTTACTAAATAATCAAAAAATATGC	200			
326	TCAAGACCCAGCGCTGGCCAACTGATGAATCCCACTCTCTACTAAATAATCAAAAAATTAAC	267			
201	TGGGCAATGGTGGCAACACCTGTGTAGTCCAGACTCTACTAGAGACCGGAGATTGCAGTAGC	260			
266	TGGGCAATGATGTGACAGCTCTGTATATCCAGACTTCTGGAGGCTGAGAGTTGCAGTAGC	207			
261	TGAGATCGCAGAGTGAGCCGAATCAAGATCAAGAGTGAGCAGAGTGAGACACCCCT	320			
206	CAAGATGCACAATCGCACTCCAGCTCGGGTGAGCAGAGCAGAGCTCGCTCCCAAAAAA	147			
321	CAAAAAACAACAAAAACAAAAA	347			
146	TAAATTAATTAATTAATTAATTAATTAATCA	120			

[illegible]

Email: li3-cdna@nifty.com
NEBO human cDNA project (New Energy and Industrial Technology Developmental Organization, Japan): cDNA library construction: Helix Research Institute (HRI): 5'-end one pass sequencing: HRI, Research Association for Biotechnology (RAB) and Biotechnology Center, National Institute of Technology and Evaluation; 3'-end one pass sequencing: RAB.

FEATURES	Location/Qualifiers
source	1. .581

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/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/clone="OCBBF3024884"
/tissue_type="brain"
/dev_stage="fetal"
/clone_lib="OCBBF3"
/notes="Vector: pME18SFL3"

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	Query Match	36.8%	Score 147	DB 9	Length 581
	Best Local Similarity	71.9%	Pred. No. 2,7e-18		
	Matches	Conservative	0	Mismatches 75	Indels 0
				Gaps	0
QY	81	ATGCTCTTAATCCCAAGCACTTGTGGGAGGCCAAGGTGGSCGGATCACTTGAGGTCAAGAGA	140		
DB	579	ACGGCTTAATTAATCCAGACATTGTGGAGGGCTGAGGCCGGTGGATCACTTGAGGTCAAGAGT	520		
QY	141	TCGAGACCATCTGTGGCAACATGTGTGAACCCCGTCTTATCTAAATAATATCAAAAAATTAGC	200		
DB	519	TCAGAACCAGGCTGGCCAAACAGTGAATACCCACTCTTACTTAAATAATATCAAAAAATTAGC	460		
QY	201	TGGGCATGTGTGGCACACCTGTGTAGTCCGAGTACTCAGAGACCGGAGATTGCACTGAGC	260		
DB	459	TGGGCATGTGTGTGCATGCTCTGTATATCCACACTTATTTGGGAGGCTGAGAGTTGCAGGTGAGC	400		
QY	261	TGAGATCGCAGAGTGAGCCGGAATACACAGATCACAGATGTGACAGAGTGAAGCKCCGCT	320		
DB	399	CAAGTGGCCACAACTGCACCTCGAGCTGGGTGACAGAGCAAGACCTCCCTCTCCAAAAAA	340		
QY	321	CAAAAACACACAAAAAACAATAAA	347		
DB	339	TAAATTAATTAATTAATTAATTAATCA	313		

RESULT	23
BM542252/c	
LOCUS	BM542252
DEFINITION	1085 bp mRNA linear EST 20-FEB-2002
	AGENCOURT_6436650 NIH_MGC_72 Homo sapiens cDNA clone IMAGE:5539645
	5', mRNA sequence.
ACCESSION	BM542252
VERSION	BM542252.1
KEYWORDS	GI:18771599
SOURCE	EST.
	Homo sapiens (human)
ORGANISM	Homo sapiens

REFERENCE	1 (bases 1 to 1085)
AUTHORS	NIH-MGC http://mgc.nci.nih.gov/ .
TITLE	National Institutes of Health, Mammalian Gene Collection (MGC)
JOURNAL	Unpublished (1999)
COMMENT	Contact: Robert Strausberg, ph.D.

Email: cgapds-remail.nih.gov
Tissue Procurement: ATCC/DCFD/STP
CDNA Library Preparation: Life Technologies, Inc.
CDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
DNA Sequencing by: Agencourt Bioscience Corporation
Clone distribution: MGC clone distribution information can be
found through the I.M.A.G.E. Consortium/LLNL at:
<http://image.llnl.gov>
Plate: LLAM12234 row: h column: 14
High quality sequence stop: 418.

FEATURES

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/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/clone="IMAGE:5539645"
/tissue_type="melanotic melanoma"
/lab_host="DH10B (phage-resistant)"
/clone_id="NTH MGC 72"
/notes="Organ: skin; Vector: pCMV-Sport6; Site_1: NotI; Site_2: SalI; Cloned unidirectionally. Primer: Oligo dT Average insert size 2 kb. Library constructed by Life technologies."

```

ORIGIN

Query Match	36.8%	Score 147	DB 2	Length 1085
Best Local Similarity	71.9%	Pred. No. 12	4e-18	
Matches 192	Conservative	0	Mismatches 75	Indels 0
Qy	81	ATGCCTTAATCCAGACACTTCGGGAGGCCAAGGTGGGGCGATCACTGAGGTCAAGAGA	140	
Db	329	ACGCCTTAATCCAGACACTTTTGGAGGCTGAGGCCGCTGATCACTGAGGTCAAGAGT	270	
Qy	141	TCGAGACCATCTGGCCACATGGTGAACCCCGCTCTTTACTAAAAATACAAAAATAGC	200	
Db	269	TCAGAGCCACGCTGGCCACACAGTGAACCCCATCTCTACTAAAAATACAAAAATTAGC	210	
Qy	201	TGGGCATGTGTGSCACACACTGTAGTCCAGCTACTCAGAGCCGGAGATTGCAGTAGC	260	
Db	209	TGGGCATGTAGTGTGCATAGCTGTAAATCCAGCTACTTTGGAGGCTGAGGTTGCAGTAGC	150	
Qy	261	TGAGATCGCAGAGTAGAGCCGAAAATCACAGATCAACAGGTAGCAGAGTGAAGACCCGCTT	320	
Db	149	CAAGATCGCACAACATGCACTCCAGCGCTGGGTGAACAGCAAGACTCGCTCCCAAAAAA	90	
Qy	321	CAAAAACACACAAAAAAACAAAAA	347	
Db	89	TAAATAAATAAATAAATAAATATACA	63	

DEFINITION RPC11-165E2.TV RPC1-11 Homo sapiens genomic clone RPC1-11-165E2, genomic survey sequence.

ACCESSION AQ381551

VERSION AQ381551.1 GI:4352574

KEYWORDS GSS.

SOURCE Homo sapiens (human)

ORGANISM Homo sapiens

REFERENCE Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini; Homnidae; Homo.

AUTHORS Zhao, S., Adams, M.D., Niernan, W., Malek, J., de Jong, P., and Venter, J. C.

TITLE Use of BAC End Sequences from Library RPC1-11 for Sequence-Ready Map Building

JOURNAL Unpublished (1997)

COMMENT Other GSSs: RPC11-165E2.TJ
Contact: Shaying Zhao, William Niernan, Mark Adams
Department of Eukaryotic Genomics
The Institute for Genomic Research
9712 Medical Center Dr., Rockville, MD 20850
Tel: 301 838 0200
Fax: 301 838 0208
Email: hbe@tigr.org
Clones are derived from the human BAC library RPC1-11. For BAC library availability, please contact Pieter de Jong (pieter@lejonc.med.buffalo.edu). Clones may be purchased from BACPAC Resources (http://bacpac.med.buffalo.edu/ordering) or from Research Genetics (info@resgen.com). BAC end search page: http://www.tigr.org/cdb/humgen/bac_end_search/bac_end_search.html
Seq primer: T7
Class: BAC ends.

FEATURES

source Location/Qualifiers

1..536

/organism="Homo sapiens"

/mol_type="genomic DNA"

/db_xref="GDB:7563073"

/db_xref="taxon:9606"

/clone="RPC1-11-165E2"

/sex="Male"

/cell_type="Lymphocytes"

/clone_id="RPC1-11"

/note="Vector: pBAC3.6; Site 1: EcoRI; Site 2: EcoRI; RPC11 Human Male BAC Library"

ORIGIN

Query Match 36.7%; Score 146.8; DB 11; Length 536;
Best Local Similarity 71.7%; Pred. No. 2.9e-18;
Matches 208; Conservative 1; Mismatches 73; Indels 8; Gaps 1;

84 CCTGTAATCCAGCACTTCGGAGGCCAAGTGGCGGATCACTGAGTCAAGAGATCG 143
|||||
441 CCTGTAATCCAGCACTTCGGAGGCCAAGTGGCGGATCACTGAGTCAAGATTTG 382
|||||
144 AGACCATCTCGGCCAATCATGTGAAACCCCGCTTTACTTAAATAACAAAAATAGCTGG 203
381 AGACCAAGCTGGCCAACTGTAAGAACCCCATTTCTATTAATAATACAAAAATTAGCTGG 322
|||||
204 GCATGTGGACACACACTGTAGTCCAGCTACTCAGAGCCGAGATTTGACAGTAGCTGA 263
321 GCGTGTGGTGGGTGCTGTATCCAGCTACTCAGAGGACAGAGTTGTAGTAGC--- 265
264 GATGGCAGATGACCCGAATACACAGATCACAAGTAGAGAGAGATGAGAACTTCATCTCAA 323
264 -----CCAAAGATTGCACCACTGCACTCTGCGCGACAGAGTGAAGACTTCATCTCAA 210
324 AAACAACAACAAAAACAAAAACATTAAGACATTGTCATCTGCGGTT 373
209 AAAAAAAGAAAAAGAAAAAGAAAAACATTAAGCTGAAATTTTAGGGGT 160

LOCUS BQ432755 775 bp mRNA linear EST 24-MAY-2002

DEFINITION AGENCOURT 7860001 NIH_MGC_55 Homo sapiens cDNA clone IMAGE:6108747 5', mRNA sequence.

ACCESSION BQ432755

VERSION BQ432755.1 GI:21171831

KEYWORDS EST.

SOURCE Homo sapiens (human)

ORGANISM Homo sapiens

REFERENCE Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini; Homnidae; Homo.

AUTHORS NIH-MGC http://mgs.nci.nih.gov/.

TITLE National Institutes of Health, Mammalian Gene Collection (MGC)

JOURNAL Unpublished (1999)

COMMENT Contact: Robert Strausberg, Ph.D.
Email: sgabs@remail.nih.gov
Tissue Procurement: ATCC
CDNA Library Preparation: CLONTECH Laboratories, Inc.
CDNA Library Arrayed by: The I.M.A.G.E. Consortium (LNL)
DNA Sequencing by: Agencourt Bioscience Corporation
Clone distribution: MGC clone distribution information can be found through the I.M.A.G.E. Consortium/LNL at:
http://image.lnl.gov
Plate: LNCM353 row: 1 column: 04
High quality sequence stop: 555.

FEATURES

source Location/Qualifiers

1..775

/organism="Homo sapiens"

/mol_type="mRNA"

/db_xref="taxon:9606"

/clone="IMAGE:6108747"

/tissue_type="from acute myelogenous leukemia"

/lab_host="DH10B (T1 phage-resistant)"

/clone_id="NIH_MGC_55"

/note="Organ: bone marrow; Vector: pDNR-LIB (Clontech); Site 1: SfiI (ggcgctcgccg); Site 2: SfiI (ggcgatcgccg); Double-stranded cDNA was prepared from cell line RNA. 5' and 3' adaptors were used in cloning as follows: 5' adaptor sequence: 5'-CACGCCATTATGCCC-3' and 3' adaptor sequence: 5'-ATTCTAGAGCCGAGGCGCCGACATG-dt(30)BN-3' (where B = A, C, or G and N = A, C, G, or T). Average insert size 1.65 kb (range 0.9-4.0 kb). 14/15 colonies contained inserts by PCR. This library was enriched for full-length clones and was constructed by Clontech Laboratories (Palo Alto, CA)."

ORIGIN

Query Match 36.7%; Score 146.8; DB 3; Length 775;
Best Local Similarity 72.5%; Pred. No. 2.7e-18;
Matches 190; Conservative 0; Mismatches 72; Indels 0; Gaps 0;

81 ATGCGTGAATCCAGCACTTCGGAGGCCAAGTGGCGGATCACTGAGTCAAGAGA 140
645 ACGCTGTAATCTCAGCACTTTGGAGGCCGAGGACAGTGGATCACTGAGTCAAGAGT 586
|||||
141 TCGGACCATCTGCGCCAACTGTGTAACCCCGCTTTACTTAAATAACAAAAATAGTC 200
585 TCAAGACTGAGCTGGCCAACTGTGTAACCCCGCTTTACTTAAATAACAAAAATAGT 526
201 TGGCATGTGGACACACACTGTAGTCCAGCTACTCAGAGCCGAGATTTGACAGTAGC 260
525 TGGCATGTGGACATGACCTGTATCCGCTACTCTCGGAGGCTGAGGTTCAGTAGC 466
261 TGAGATCGCAGATGAGCCGAATACAGATCACAAGTAGAGAGAGTGAAGACCCGCTCT 320
465 CAAGATTGACCACTGCAACCCCACTGGGCAAAACAGTAGACTTTTCTCAAAAAAAA 406
321 CAACAACAACAACAAAAACA 342
405 AAAAAATTAATATATAGAAAA 384

RESULT 26
 A1753536/c 469 bp mRNA linear EST 20-JUN-2002
 LOCUS cr11h09.x1 Human bone marrow stromal cells Homo sapiens CDNA clone
 DEFINITION HBMS.Cr11h09 3', mRNA sequence.
 ACCESSION A1753536
 VERSION A1753536.1 GI:5131800
 KEYWORDS EST.
 SOURCE Homo sapiens (human)
 ORGANISM Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini;
 Homidae; Homo.
 1 (bases 1 to 469)
 Jia, L., Young, M.F., Powell, J., Yang, L., Ho, N.C., Hotchkiss, R.,
 Robey, P.G. and Francomano, C.A.
 Gene expression profile of human bone marrow stromal cells:
 high-throughput expressed sequence tag sequencing analysis
 Genomics 79 (1), 7-17 (2002)
 11827452
 COMMENT Contact: Libin Jia
 Medical Genetics Branch
 National Human Genome Research Institute
 10/10C101, 9000 Rockville Pike, Bethesda, MD 20892-1267, USA
 Tel: 301-402-4877
 Fax: 301-496-7157
 Email: libin@nih.gov
 DNA Sequencing and analyses by National Institutes of Health
 Intramural Sequencing Center (NISC).
 Plate: 11 row: h column: 09
 Seq primer: -21M3 forward primer (ABI).
 Location/Qualifiers
 1..469
 /organism="Homo sapiens"
 /mol_type="mRNA"
 /db_xref="taxon:9606"
 /clone="HBMSC_cr11h09"
 /sex="mixed"
 /tissue_type="bone marrow stroma"
 /dev_stage="mixed"
 /lab_host="XLI-Blue MRF/SOLR"
 /clone_lib="Human bone marrow stromal cells"
 /note="Vector: pBluescript; Site_1: EcoRI; Site_2: XhoI;
 mRNA made from human bone marrow stroma, CDNA made by
 oligo-dT priming. Directionally cloned. Size-selected for
 average insert size >0.5 kb. Library constructed by Dr.
 Marian Young and Dr. Pamela Gehron Robey (NIDCR). Library
 supplied by Dr. Libin Jia (NHGRI)"

ORIGIN
 Query Match 36.7%; Score 146.6; DB 1; Length 469;
 Best Local Similarity 67.9%; Pred. No. 3.3e-18;
 Matches 218; Conservative 1; Mismatches 94; Indels 8; Gaps 1;

33 ACCCAATTTAATTAAGACTTTGTCAGGCCGAGCATGACCTGCTGATGCTTATATC 92
 |||
 321 ACNCAATTTTACATTAAGAAATACCTGTGACAGCCATGCTGTGATCAGAGCTTGTATTC 262
 |||
 93 CCAGCACTTCGGAGAGCCGAAGTGGGCGGATCACTGAGTCAAGATCGAGACCATCC 152
 |||
 261 CCAGCACTTTGGAGAGCCGAAGTGGGCGGATCACTGAGTCAAGATCGAGACCATCC 202
 |||
 153 TGGCCACATGAGTGAACCCCGTCTTTTACTTAAATAACAAAATAGCTGGGCGATGTGG 212
 |||
 201 TTGCCACATATAGTGAACCCGTCTCTACTAAATAACAAAATAGCTGGGCGATGTGG 142
 |||
 213 CACACACTGTATGTCAGCTTACTCAAGAG-----CCGAGATTGCGAGTGAAGTGAAG 264
 |||
 141 CAGGCACTGTATGTCAGCTTACTCAAGAGAGGCTTTTGAACCCGAGAGCAGAGGTTGAG 82
 |||
 265 ATCCGAGATGAGCCGAATATCAAGATCAAGAGTGAAGCAGAGTGAAGACCGCTCTCAAA 324
 |||

Db 81 CGAGCTGAGATGCGCGCACTGCATCCAGCANTGGGTGATAGAGTGAATTCACTTCCAA 22
 QY 325 AACCAACACACAAAAACAAAA 345
 |||
 Db 21 AAAAAAAAAAAAAAAAAAAAAA 1

RESULT 27
 A1434037/c 490 bp mRNA linear EST 30-MAR-1999
 LOCUS t13b10.x1 NCI CGAP Lym12 Homo sapiens CDNA clone IMAGE:2132539 3'
 DEFINITION similar to contatns Alu repetitive element; mRNA sequence.
 ACCESSION A1434037
 VERSION A1434037.1 GI:4293374
 KEYWORDS EST.
 SOURCE Homo sapiens (human)
 ORGANISM Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini;
 Homidae; Homo.
 1 (bases 1 to 490)
 NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
 National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
 Tumor Gene Index
 Unpublished (1997)
 COMMENT Contact: Robert Strausberg, Ph.D.
 Email: cgapdb-remail.nih.gov
 Life Technologies catalog #: 11547-015
 DNA Sequencing by: Washington University Genome Sequencing Center
 Clone distribution: NCI-CGAP clone distribution information can be
 found through the I.M.A.G.E. Consortium/BLN at:
 www.bio.linn.gov/bdtp/image/image.html
 Insert Length: 1365 Std Error: 0.00
 Seq primer: -40UP from Gldco
 High quality sequence stop: 297.
 Location/Qualifiers
 1..490
 /organism="Homo sapiens"
 /mol_type="mRNA"
 /db_xref="taxon:9606"
 /clone="IMAGE:2132539"
 /tissue_type="lymphoma, follicular mixed small and large
 cell"
 /lab_host="DH10B"
 /clone_lib="NCI CGAP Lym12"
 /note="Organ: lymph node; Vector: pCMV-SPORT6; Site_1:
 SalI; Site_2: NotI; Cloned unidirectionally. Primer:
 Oligo dT. Average insert size 1.25 kb. Life Technologies
 catalog #: 11547-015"

ORIGIN
 Query Match 36.7%; Score 146.6; DB 1; Length 490;
 Best Local Similarity 75.5%; Pred. No. 3.3e-18;
 Matches 182; Conservative 0; Mismatches 59; Indels 0; Gaps 0;

63 AGCATGACATCTGGCTGATGCTGTATTCACAGCACTTCGGAGGCCAAGTGGCGGA 122
 |||
 285 AGGCAGAGCAGATGCTCATCTCTGTATCCAGACATTTGGAGGCCAAGTGGCGGA 226
 |||
 123 TCACCTGAGTCAAGAGATGAGACCATCTGGGCCAATGATGTGAACCCGCTTTTACT 182
 |||
 225 TCATCTGAGTCAAGAGATGAGACCATCTGGGCCAATGATGTGAACCCGCTTTTACT 166
 |||
 183 AAAAAATCAAAAAATAGCTGGGCGATGTGGCACAACCTGTAGTCCAGTACTCAGAG 242
 |||
 165 AAAAAATCAAAAAATAGCTGGGCGATGTGGCACAACCTGTAGTCCAGTACTCAGAG 106
 |||
 243 CCGAGATTGAGTGAAGTGAAGTGAAGTGAAGTGAAGTGAAGTGAAGTGAAGTGAAG 302
 |||
 105 GCTAGGATGAAGATATCACTTGAACCCGAGGAGGAGGCTTCAGTGAAGCGAGATTACG 46
 |||
 303 C 303

Db 45 C 45

RESULT 28
AG156377/c
LOCUS
DEFINITION AG156377 667 bp DNA linear GSS 09-JAN-2002
Pan troglodytes DNA, clone: RP43-020P17.T7, genomic survey

ACCESSION AG156377
VERSION AG156377
KEYWORDS GI:16686055
SOURCE GSS.
ORGANISM Pan troglodytes (chimpanzee)
Pan troglodytes
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini;
Homnidae; Pan.

REFERENCE 1
AUTHORS Fujiyama, A., Hattori, M., Toyoda, A., Taylor, T.D., Yada, T.,
Toroki, Y., Matanabe, H. and Sakaki, Y.
TITLE BAC end sequences of library RP43-020P17.T7
JOURNAL Unpublished
REFERENCE 2 (bases 1 to 667)
AUTHORS Fujiyama, A., Hattori, M., Toyoda, A., Taylor, T.D., Yada, T.,
Toroki, Y., Matanabe, H. and Sakaki, Y.
TITLE Direct Submission
JOURNAL Submitted (02-AUG-2001) Ageo Fujiyama, The Institute of Physical
and Chemical Research (RIKEN), Genomic Sciences Center (GSC);
1-7-22 Suehiro-cho, Tsurumi-ku, Yokohama, Kanagawa 230-0045, Japan
(E-mail: chimpanzee@sc.riken.go.jp, URL: http://hgp.gsc.riken.go.jp/,
Tel: 81-45-503-9111, Fax: 81-45-503-9170)
COMMENT Clones are derived from the chimpanzee BAC library RP43-020P17.T7. This BAC
end was generated during the R&D process and may have higher chance
of clone tracking errors.
PRIMERS

Sequencing: T7
LIBRARY
Vector : pBACe3.6
R Site 1 : EcoRI
R Site 2 : EcoRI
Location/Qualifiers
1. 667
/organism="Pan troglodytes"
/mol_type="genomic DNA"
/db_xref="taxon:9598"
/clone="RP43-020P17.T7"
/sex="male"
/cell_type="lymphocytes"
/clone_lib="RP43-020P17.T7 Chimpanzee Male BAC library"

ORIGIN
Query Match 36.7%; Score 146.6; DB 14; Length 667;
Best Local Similarity 78.2%; Pred. No. 3.1e-18;
Matches 176; Conservative 0; Mismatches 49; Indels 0; Gaps 0;

FEATURES
source
1. 667
/organism="Pan troglodytes"
/mol_type="genomic DNA"
/db_xref="taxon:9598"
/clone="RP43-020P17.T7"
/sex="male"
/cell_type="lymphocytes"
/clone_lib="RP43-020P17.T7 Chimpanzee Male BAC library"

ORIGIN
Query Match 36.7%; Score 146.6; DB 14; Length 667;
Best Local Similarity 78.2%; Pred. No. 3.1e-18;
Matches 176; Conservative 0; Mismatches 49; Indels 0; Gaps 0;

Db 81 ATGCTGTAATCCGACGACTTCGGAGGCGAAGTGGGGGATCACTGAGTCAAGAGA 140
Db 559 ATGCTGTAATCCGACGACTTCGGAGGCGAAGTGGGGGATCACTGAGTCAAGAGA 500
Qy 141 TCGAGACCACTCTGGCCCAATGTGTAACCCCGCTTTTACTTAAATAAATAATAGC 200
Db 499 TTGAGACCACTCTGGCCCAATGTGTAACCCCGCTTTTACTTAAATAAATAATAGC 440
Qy 201 TGGGCGATGTTGGCACACACCTGTAGTCCAGGTAATCAAGAGCCGAGATTGCAATGAGC 260
Db 439 TGGGCGATGTTGGCACACACCTGTAGTCCAGGTAATCAAGAGCCGAGATTGCAATGAGC 380
Qy 261 TGAGATCGAGAGTGGAGCGAATATCAAGATCAAGAGTGGAGC 305
Db 379 CTGGAACCTGGAGGCGAGTTTGCAGTGAAGATTGAGCCG 335

LOCUS
DEFINITION
ACCESSION
VERSION
KEYWORDS
SOURCE
ORGANISM

AW902341 406 bp mRNA linear EST 24-MAY-2000
OV3-NN1023-130500-179-b08 NN1023 Homo sapiens cDNA, mRNA sequence.
AW902341 GI:8066546
EST.
Homo sapiens (human)
Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini;

REFERENCE
AUTHORS
1 (bases 1 to 406)
Dias Neto, E., Garcia Correa, R., Verjovski-Almeida, S., Brienes, M.R.,
Nagai, M.A., da Silva, W. Jr., Zago, M.A., Bordin, S., Costa, F.F.,
Goldman, G.H., Carvalho, A.F., Matsukuma, A., Bala, G.S., Simpson, D.H.,
Brunstein, A., de Oliveira, P.S., Bucher, P., Jongeneel, C.V.,
O'Hare, M.J., Soares, F., Brentani, R.R., Reis, L.F., de Souza, S.J. and
Simpson, A.J.
TITLE Shotgun sequencing of the human transcriptome with ORF expressed
sequence tags
JOURNAL Proc. Natl. Acad. Sci. U.S.A. 97 (7), 3491-3496 (2000)
PUBMED 10737800
COMMENT Contact: Simpson A.J.G.
Laboratory of Cancer Genetics
Ludwig Institute for Cancer Research
Rua Prof. Antonio Prudente 109, 4 andar, 01509-010, Sao Paulo-SP,
Brazil
Tel: +55-11-2704922
Fax: +55-11-2707001
Email: asimpson@ludwig.org.br
This sequence was derived from the FAPESP/LICR Human Cancer Genome
Project. This entry can be seen in the following URL
(http://www.ludwig.org.br/scripts/gethtml2.pl?tl=et2-QV3-NN1023-130
500-179-b08&et3=2000-05-13&t4=1)
Seq primer: puc 18 forward
High quality sequence start: 22
High quality sequence stop: 406.
Location/Qualifiers
1. 406
/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/dev_stage="Adult"
/clone_lib="NN1023"
/note="Organ: nervous normal; Vector: puc18; Site 1: SmaI;
Site 2: SmaI; A mini-library was made by cloning products
derived from ORSITES PCR (U.S. Letters Patent application
No. 196,716 - Ludwig Institute for Cancer Research)
profiles into the puc 18 vector. Reverse transcription of
tissue mRNA and cDNA amplification were performed under
low stringency conditions."

ORIGIN
Query Match 36.6%; Score 146.4; DB 7; Length 406;
Best Local Similarity 86.2%; Pred. No. 3.7e-18;
Matches 162; Conservative 0; Mismatches 26; Indels 0; Gaps 0;

Qy 81 ATGCTGTAATCCGACGACTTCGGAGGCGAAGTGGGGGATCACTGAGTCAAGAGA 140
Db 301 AGCGCTGTAATCCGACGACTTCGGAGGCGAAGTGGGGGATCACTGAGTCAAGAGA 242
Qy 141 TCGAGACCACTCTGGCCCAATGTGTAACCCCGCTTTTACTTAAATAAATAATAGC 200
Db 241 TCGAGACCACTCTGGCCCAATGTGTAACCCCGCTTTTACTTAAATAAATAATAGC 182
Qy 201 TGGGCGATGTTGGCACACACCTGTAGTCCAGGTAATCAAGAGCCGAGATTGCAATGAGC 260
Db 181 TGGGCGATGTTGGCACACACCTGTAGTCCAGGTAATCAAGAGCCGAGATTGCAATGAGC 122
Qy 261 TGAGATCG 268
Db 121 CGAGATAG 114

RESULT 30
 A1302156/c 524 bp mRNA linear EST 03-DEC-1998
 LOCUS qn58a05.x1 NCI_CGAP_Kids Homo sapiens cDNA clone IMAGE:1902416 3
 DEFINITION similar to contigins Alu repetitive element, mRNA sequence.
 ACCESSION A1302156
 VERSION A1302156.1 GI:3961502
 KEYWORDS EST.
 SOURCE Homo sapiens (human)
 ORGANISM Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini;
 Hominidae; Homo.
 REFERENCE 1 (bases 1 to 524)
 NCI-CGAP <http://www.ncbi.nlm.nih.gov/ncicgap>.
 TITLE National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
 Tumor Gene Index
 JOURNAL Unpublished (1997)
 COMMENT Contact: Robert Strausberg, Ph.D.
 Email: cgapbs-remail.nih.gov
 Tissue Procurement: Christopher Moskalko, M.D., Ph.D., Michael R.
 Emert-Buck, M.D., Ph.D.
 cDNA Library Preparation: M. Bento Soares, Ph.D.
 DNA Library Arrayed by: Greg Lennon, Ph.D.
 DNA Sequencing by: Washington University Genome Sequencing Center
 Clone distribution: NCI-CGAP clone distribution information can be
 found through the I.M.A.G.E. Consortium/LLNL at:
www-bio.llnl.gov/bbrp/image/image.html
 Seq primer: -40UP from G1bco
 High quality sequence stop: 461.
 Location/Qualifiers
 1..524
 /organism="Homo sapiens"
 /mol_type="mRNA"
 /db_xref="taxon:9606"
 /clone="IMAGE:1902416"
 /cissue_type="2 pooled tumors (clear cell type)"
 /lab_host="DH10B"
 /clone_lib="NCI_CGAP_Kids"
 /note="Organ: Kidney; Vector: pT7T3D-PacI; Site_1: Not I;
 Site_2: Eco RI; 1st strand cDNA was primed with a Not I -
 oligo (dT) primer [5',
 AACTGGAAGAATTCGGCGCCGCAATATTTTTTTTTTTTTTTT 3'],
 double-stranded cDNA was ligated to Eco RI adaptors
 (Pharmacia), digested with Not I and cloned into the Not I
 and Eco RI sites of the modified pT7T3 vector. Library
 went through one round of normalization. Library
 constructed by Bento Soares and M. Patricia Bonaldo."

ORIGIN
 Query Match 36.6%; Score 146.4; DB 1; Length 524;
 Best Local Similarity 72.6%; Pred. No. 3.5e-18;
 Matches 209; Conservative 1; Mismatches 62; Indels 16; Gaps 1;

81 ATGCTGTAATCCCGACGACTTCGGGAGCGCAAGTGGGGGATCACTGAGGTCAAGA 140
 450 ATGCTGTAATCCCGACGACTTCGGGAGCGCGGGGATCACTGAGGTCAAGA 391
 141 TCGAGACCATCTCTGCGCAACATGTGAAACCCCGCTTACTAATAAATACAAAATATAGC 200
 390 TCGAGACCATCTGCGCAACATGTGAAACCCCGCTTACTAATAAATACAAAATATAGC 331
 201 TGGGATATGTCACACACCTGTAGTCCAGCTACTCAGAGACCGGAGATTGCACTGAGC 260
 330 TGGGATATGTCACACACCTGTAGTCCAGCTACTCAGAGATTGAGAGAGTGG 271
 261 TGAAGTGCAGAGTGAAGCGGAATATCAAGATCAAGAGTGAAG-----CA 304
 270 CTGAGCTCTGGAGGTGAAGTGAAGCGGAGACCGGTCTCCAGACCTGGGTACA 211
 305 GAGTGAAGACCGCTCTCAAAAACAACAACAAAACAAAACAAAACATA 352
 210 GAGCGAGACTCTGTCTCAAAAATATATATATATATATATATATATAA 163

RESULT 31
 CR860050 1459 bp mRNA linear HTC 12-NOV-2004
 LOCUS Pongo pygmaeus mRNA; cDNA DKFZp470B1412 (from clone DKFZp470B1412).
 DEFINITION
 ACCESSION CR860050
 VERSION CR860050.1 GI:55730961
 KEYWORDS HTC.
 SOURCE Pongo pygmaeus (orangutan)
 ORGANISM Pongo pygmaeus
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini;
 Hominidae; Pongo.
 REFERENCE 1 (bases 1 to 1459)
 Wamburt, R., Heubner, D., Mewes, H.W., Weil, B., Anld, C., Osanger, A.,
 Fodor, G., Han, M. and Wiemann, S.
 TITLE The German cDNA Consortium
 JOURNAL Submitted (12-NOV-2004) MIPS, Ingolstaedter Landstr.1, D-85764
 Neuenberg, GERMANY
 COMMENT Clone from S. Wiemann, Molecular Genome Analysis, German Cancer
 Research Center (DKFZ); Email s.wiemann@dkfz-heidelberg.de;
 sequenced by Agowa (Berlin/Germany) within the cDNA sequencing
 consortium of the German Genome Project.
 This clone (DKFZp470B1412) is available at the RZPD Deutsches
 Ressourcenzentrum fuer Genomforschung GmbH in Berlin, Germany.
 Please contact RZPD for ordering:
<http://www.rzpd.de/cgi-bin/products/cl.cgi?cloneID=DKFZp470B1412>
 Further information about the clone and the sequencing project is
 available at <http://mips.gsf.de/projects/cdna/>.

FEATURES

source

1..1459
 Location/Qualifiers
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 /organism="Pongo pygmaeus"
 /mol_type="mRNA"
 /db_xref="taxon:9600"
 /clone="DKFZp470B1412"
 /cissue_type="liver"
 /clone_lib="470 (synonym: pliv1). Vector pSport1_Sfi; host
 DH10B; sites SfiI + SfiIb"
 /dev_stage="adult"
 /note="NGF-beta receptor interacting protein 1 (Homo
 sapiens)"
 1..1459
 /gene="DKFZp470B1412"
 15..992
 /gene="DKFZp470B1412"
 /codon_start=1
 /product="hypothetical protein"
 /protein_id="CAH92198.1"
 /db_xref="GI:55730962"
 /translation="MKPILQGHRSITQIKYRNEGDLFTVAKDPIVNVVYVNGER
 LGTYMGHTGAWCVADMDTGTGVLTSADNSCLMDCTGALALKTNSAVRTGFP
 FGSIIMSTTDKMGYOCFVSPFLRDSOI DNNBPWKIKCNSTKITSAYWPGIQC
 IIAHBSGLNLYSAKSEVYVYVKESRQINDIQLSDMTPTTASQNTAKLPDST
 TLHOKTRFIRPNVNSALSPNYHVLGSGQEMADVTTSTRIGKFAKRFPHLAFEE
 EFGVKGHFGLINSVAHPHPDKSYSSGDEGVRIHPDPYFEPFEBA"

ORIGIN

Query Match 36.6%; Score 146.4; DB 6; Length 1459;
 Best Local Similarity 73.8%; Pred. No. 2.9e-18;
 Matches 186; Conservative 0; Mismatches 66; Indels 0; Gaps 0;

46 TAAAGCATTTGTCAAGCCAGCATGCACTGGCTGAATCCGTATATCCAGACCTTCCGG 105
 990 TAAAGAGTGTGATCTCTGCGCAGGCGGGGTGCTCATGCTGTATATCCACACACTTTGG 1049
 106 AGGCCAAGTGGGGGATCACTGAGGTCAAGAGATGAGACCATCTGCGCAACATGCT 165
 1050 AGGCCAAGTGGGGGATCACTGAGGTCAAGAGATTGAGACCAAGCTGCAACATGGA 1109
 166 GAAACCCCGCTTACTAATAAATCAAAAATAGCTGGGATGAGTGGGACACACTGTAG 225

Db 1110 GAAACCCGCTCTACTATAAATAAATAATAGCCGGGAGGATGGGACACGCGTATAG 1169
 QY 226 TCCGAGCTACTCAGAGCGCGGAGATTGCGAGTGAGTCGACAGAGTACCGGAATC 285
 Db 1170 TCCGAGCTACTCAGAGCGGAGGCTGAGGACGAGATCATCTTGAACCCAGAGGAGAGGTTGC 1229
 QY 286 ACAGATCAGACA 297
 Db 1230 AGTAGCTGAGA 1241

RESULT 32
 LOCUS CN268905 302 bp mRNA linear EST 16-MAY-2004
 DEFINITION 17000597786250 GRN PREHBP Homo sapiens CDNA 5', mRNA sequence.
 ACCESSION CN268905
 VERSION CN268905.1 GI:47285319
 KEYWORDS EST.
 SOURCE Homo sapiens (human)
 ORGANISM Homo sapiens

REFERENCE
 AUTHORS Brandenberger, R., Wei, H., Zhang, S., Lei, S., Murage, J., Fisk, G.J., Li, Y., Xu, C., Fang, R., Guegler, K., Rao, M.S., Mandalam, R., Lebkowski, J. and Stanton, L.W.
 TITLE Transcriptional characterization elucidates signaling networks that control human ES cell growth and differentiation
 JOURNAL Nat. Biotechnol. 22 (6), 707-716 (2004)
 PUBMED 15146197
 COMMENT Contact: Brandenberger R
 Regenerative Medicine
 Geron Corporation
 230 Constitution Drive, Menlo Park, CA 94025, USA
 Tel: 650 473 8658
 Fax: 650 473 7760
 Email: rbrandenberger@geron.com
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FEATURES
 Location/Qualifiers

1..302
 /organism="Homo sapiens"
 /mol_type="mRNA"
 /db_xref="taxon:9606"
 /tissue_type="embryonic stem cells, DMSO-treated H9 cell line"
 /clone_lib="GRN_PREHBP"
 /note="Oligo dt primed, full-length enriched cDNA library from DMSO-treated hES cell line H9 (p22) maintained in feeder-free conditions"

ORIGIN
 Query Match 36.6%; Score 146.2; DB 8; Length 302;
 Best Local Similarity 78.5%; Pred. No. 4,3e-18;
 Matches 175; Conservative 0; Mismatches 48; Indels 0; Gaps 0;

QY 81 ATGCTGTATATCCGACGACTTGGGAGCGCAAGTGGCGGATCACTGAGTCAAGA 140
 Db 299 ACGCTGTATATCCGACGACTTGGGAGCGCAAGTGGCGGATCACTGAGTCAAGA 240
 QY 141 TCGAGACCATCTCTGGCCCAATGATGTAACCCCGTCTTACTATAAATAAATAATAGC 200
 Db 239 TCAAGACCATCTCTGGCCCAATGATGTAACCCCGTCTTACTATAAATAAATAATAGC 180
 QY 201 TGGGAGTGTGGCAGACACTGTATGCTCCAGCTACTCAGAGCCGGAATTGCACTGAGC 260
 Db 179 CGGGGTGTGGCGGGCGCTGTAGTCCAGCTACTCAGAGGAGCTGAGGCAAGAAATGG 120
 QY 261 TGAGATCCGACAGTGAAGCGGAATATCAAGATACAGAGTGAAGC 303
 Db 119 CGTGAACCCGGAGGCGGAGCTTGCAGTGAAGCGGAGATTGGCG 77

RESULT 33
 LOCUS BM712012 419 bp mRNA linear EST 28-FEB-2002
 DEFINITION U1-E-DWI-ahc-a-08-0-UI.r1 U1-E-DWI Homo sapiens cDNA clone
 VERSION U1-E-DWI-ahc-a-08-0-UI 5', mRNA sequence.
 ACCESSION BM712012
 VERSION BM712012.1 GI:19025270
 KEYWORDS EST.
 SOURCE Homo sapiens (human)
 ORGANISM Homo sapiens

REFERENCE
 AUTHORS Bonaldo, M.F., Lennon, G. and Soares, M.B.
 TITLE Normalization and subtraction: two approaches to facilitate gene discovery
 JOURNAL Genome Res. 6 (9), 791-806 (1996)
 PUBMED 8889548
 COMMENT Contact: Soares, MB
 Coordinated Laboratory for Computational Genomics
 University of Iowa
 375 Newton Road, 4156 MEBRF, Iowa City, IA 52242, USA
 Tel: 319 335 8250
 Fax: 319 335 9565
 Email: bento-soares@uiowa.edu
 Tissue Procurement: Dr. Gregg Hageman
 cDNA Library preparation: Dr. M. Bento Soares, University of Iowa
 cDNA Library Arrayed by: Dr. M. Bento Soares, University of Iowa
 DNA Sequencing by: Dr. M. Bento Soares, University of Iowa
 Clone Distribution: Researchers may obtain clones from Research Genetics (www.resgen.com).

The following repetitive elements were found in this cDNA sequence: 1-181, >ALU (matched complement) 55-316, >ALU (matched complement)
 Seq primer: M13 Reverse.
 Location/Qualifiers

1..419
 /organism="Homo sapiens"
 /mol_type="mRNA"
 /db_xref="taxon:9606"
 /clone="U1-E-DWI-ahc-a-08-0-UI"
 /tissue_type="lens"
 /dev_stage="adult"
 /lab_host="DH10B (Life Technologies) (T1 phage resistant)"
 /clone_lib="U1-E-DWI"
 /note="Organ: eye; Vector: pT73-Pac (Pharmacia) with a modified polylinker; Site 1: EcoR I; Site 2: Not I; U1-E-DWI is a normalized cDNA library containing the following tissue(s): lens. The library was constructed according to Bonaldo, Lennon and Soares, Genome Research, 6:791-806, 1996. First strand cDNA synthesis was primed with an oligo-dT primer containing a Not I site. Double stranded cDNA was ligated to an EcoR I adaptor, digested with Not I, and cloned directionally into pT73-Pac vector. The oligonucleotide used to prime the synthesis of first-strand cDNA contains a library tag sequence that is located between the Not I site and the (dfl8) tail. The sequence tag for this library is CGATTAGCGA. This library was created for the program, Gene Discovery in the Visual System, supported by National Eye Institute (NEI)."

ORIGIN
 Query Match 36.6%; Score 146.2; DB 3; Length 419;
 Best Local Similarity 75.3%; Pred. No. 4e-18;
 Matches 195; Conservative 1; Mismatches 59; Indels 4; Gaps 1;

QY 81 ATGCTGTATATCCGACGACTTGGGAGCGCAAGTGGCGGATCACTGAGTCAAGA 140
 Db 299 ACGCTGTATATCCGACGACTTGGGAGCGCAAGTGGCGGATCACTGAGTCAAGA 240
 QY 141 TCGAGACCATCTCTGGCCCAATGATGTAACCCCGTCTTACTATAAATAAATAATAGC 200

DB 239 TCGAGACGAGCTGGCCCAACATGATGTAATCTCATCTACTATAAATAACAGAAATTAGC 180

QY 201 TGGGATGTTGGCACAACACTTATGCTCCAGCTACTCAGAGCCGAGATTGCACTGAGC 260

DB 179 TGGGATGTTGGTGTGATGCTGCTTAATCCAGCTACTAGAGGAGCTGAGCAGAGATCA 120

QY 261 TGAAGATCGAGAGTGAAGCCGAATATCAGATCAAGAGTGAAGAGTGAAGCKCGTCT 320

DB 119 CTGGAACCCAGAAAGTGAAGGCGACAGTAC---AGCGCCAGAGTGAAGCTCTGCT 64

QY 321 CAAAAACAACAACAAAAA 339

DB 63 CAAAAAGAAAAAGAAAAA 45

RESULT 34
CN351916 680 bp mRNA linear EST 16-MAY-2004
LOCUS CN351916.1
DEFINITION 17000532267149 GRN_EB Homo sapiens cDNA 5', mRNA sequence.
ACCESSION CN351916
VERSION CN351916.1 GI:47351850
KEYWORDS EST.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini; Homiidae; Homo.
REFERENCE 1 (bases 1 to 680)
AUTHORS Brandenberger, R., Wei, H., Zhang, S., Lei, S., Muraie, J., Flisk, G.J., Li, Y., Xu, C., Pang, R., Guegler, K., Rao, M.S., Mandalam, R., Lebowicki, J. and Stanton, L.W.
TITLE Transcriptome characterization elucidates signaling networks that control human ES cell growth and differentiation
JOURNAL Nat. Biotechnol. 22 (6), 707-716 (2004)
COMMENT 15146197
CONTACT: Brandenberger R
Regenerative Medicine
Geron Corporation
230 Constitution Drive, Menlo Park, CA 94025, USA
Tel: 650 473 8658
Fax: 650 473 7760
Email: rbrandenberger@geron.com
Insert Length: 680 Std Error: 0.00.
FEATURES
Source location/Qualifiers
1..680
/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/tissue_type="embryonic stem cells, embryoid bodies derived from H1, H7 and H9 cells"
/clone_lib="GRN_EB"
/note="Oligo dt primed, full-length enriched cDNA library from embryoid body outgrowths derived from hES cell lines H1 (p32), H7 (p29), and H9 (p26) maintained in feeder-free conditions."

ORIGIN
Query Match 36.6%; Score 146.2; DB 8; Length 680;
Best Local Similarity 89.7%; Pred. No. 3.7e-18;
Matches 157; Conservative 0; Mismatches 18; Indels 0; Gaps 0;

RESULT 35
BE379282 685 bp mRNA linear EST 21-JUL-2000
LOCUS BE379282/C 601237929P1 NIH_MGC_44 Homo sapiens cDNA clone IMAGE:3610063 5',
DEFINITION mRNA sequence.
ACCESSION BE379282
VERSION BE379282.1 GI:9324647
KEYWORDS EST.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini; Homiidae; Homo.
REFERENCE 1 (bases 1 to 685)
AUTHORS NIH-MGC http://mgi.nci.nih.gov/.
TITLE NIH-MGC National Institutes of Health, Mammalian Gene Collection (MGC)
JOURNAL Unpublished (1999)
COMMENT Contact: Robert Strusberg, Ph.D.
Email: cga@b-remail.nih.gov
Tissue Procurement: ATCC
CDNA Library Preparation: Ling Hong/Rubin Laboratory
CDNA Library Arrayed by: The I.M.A.G.E. Consortium (LNL)
DNA Sequencing by: Incyte Genomics, Inc.
Clone distribution: MGC clone distribution information can be found through the I.M.A.G.E. Consortium/LNL at:
http://image.lnl.gov
Plate: LNCM265 row: 1 column: 08
High quality sequence start: 4
High quality sequence stop: 634.
FEATURES
Source location/Qualifiers
1..685
/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/clone="IMAGE:3610063"
/tissue_type="endometrium, adenocarcinoma cell line"
/lab_host="DH10B (phage-resistant)"
/clone_lib="NIH MGC_44"
/note="Organ: uterus; Vector: pOTB7; Site 1: XhoI; Site 2: EcoRI; cDNA made by oligo-dt priming. Directionally cloned into EcoRI/XhoI sites using the following 5' adaptor: GGCACGAG(G). Library constructed by Ling Hong in the laboratory of Gerald M. Rubin (University of California, Berkeley) using ZAP-CDNA synthesis kit (Stratagene) and Superscript II RT (Life Technologies)."

ORIGIN
Query Match 36.6%; Score 146.2; DB 7; Length 685;
Best Local Similarity 75.3%; Pred. No. 3.7e-18;
Matches 195; Conservative 1; Mismatches 59; Indels 4; Gaps 1;

RESULT 36
 CN411940/c 805 bp mRNA linear EST 16-MAY-2004
 LOCUS CN411940
 DEFINITION 17000532275949 GRN ES Homo sapiens cDNA 5', mRNA sequence.
 ACCESSION CN411940
 VERSION CN411940.1 GI:47399534
 KEYWORDS EST.
 SOURCE Homo sapiens (human)
 ORGANISM Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini;
 Homiidae; Homo.
 REFERENCE 1 (bases 1 to 805)
 AUTHORS Brandenberger, R., Wei, H., Zhang, S., Lei, S., Murage, J., Fisk, G.J.,
 Li, Y., Xu, C., Fang, R., Guejler, K., Rao, M.S., Mandalam, R.,
 Ledkowski, J. and Stanton, L.M.
 TITLE Transcriptome characterization elucidates signaling networks that
 control human ES cell growth and differentiation
 JOURNAL Nat. Biotechnol. 22 (6), 707-716 (2004)
 PUBMED 15146197
 COMMENT Contact: Brandenberger R
 Regenerative Medicine
 Genon Corporation
 230 Constitution Drive, Menlo Park, CA 94025, USA
 Tel: 650 473 8658
 Fax: 650 473 7760
 Email: rbrandenberger@genon.com
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 FEATURES
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 1..805
 /organism="Homo sapiens"
 /mol_type="mRNA"
 /db_xref="taxon:9606"
 /tissue_type="embryonic stem cells, cell lines H1, H7, and
 H9"
 /clone_lib="GRN ES"
 /note="oligo dt primed, full-length enriched cDNA library
 from undifferentiated hES cell lines H1 (p32), H7 (p29),
 and H9 (p26) maintained in feeder-free conditions"
 ORIGIN
 Query Match 36.6%; Score 146.2; DB 8; Length 805;
 Best Local Similarity 75.3%; Pred. No. 3.6e-18;
 Matches 195; Conservative 1; Mismatches 59; Indels 4; Gaps 1;
 Db 81 ATGCTGTATATCCGACGACTTGGGAGGCCGAGGATCAGTCAAGTCAAGA 140
 784 AGCCTGTATATCTTGACACTTGGAGGCGGAGGATCAGTCAAGTCAAGTCAAGT 725
 QY 141 TCGAGACCATCTGGGCAACATGTGAACCCCGTCTTACTTAAATAACAAAATATAGC 200
 Db 724 TCGAAGACAGCTTGCGCAACATGTGAACCTCAATCTTCAATAAAATACAAAATTAGC 665
 QY 201 TGGGATGTGTGACACACCTGTAGTCCAGCTACTCAGAGCCGAGATTTCAGTAGC 260
 Db 664 TGGGATGTGTGATGCTGTGAATCCAGCTACTAGGAGGCTGAGGACGAGATCA 605
 QY 261 TGAAGTGGAGAGTGAAGCCGAATTCACAGATCAGAGTGAAGTGAAGTCAAGT 320
 Db 604 CTGTGAACCCAGAGGTGAAGGCGACAGTGACC---AGGCGCCAGAGTGAAGTCTGTCT 549
 QY 321 CAAAAACAACACAAAAA 339
 Db 548 CAAAAAGAAAAAAGAAAAA 530
 RESULT 37
 AA836548/c 323 bp mRNA linear EST 24-FEB-1998
 LOCUS AA836548
 DEFINITION o3dsh04.s1 NCI_CGAP Homo sapiens cDNA clone IMAGE:1370071
 similar to contains Alu repetitive element; contains element PTRS
 repetitive element ;, mRNA sequence.
 ACCESSION AA836548

VERSION AA836548.1 GI:2910867
 KEYWORDS EST.
 SOURCE Homo sapiens (human)
 ORGANISM Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini;
 Homiidae; Homo.
 REFERENCE 1 (bases 1 to 323)
 AUTHORS NCI-CGAP <http://www.ncbi.nlm.nih.gov/ncicgap>.
 TITLE National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
 Tumor Gene Index
 JOURNAL Unpublished (1997)
 COMMENT Contact: Robert Strausberg, Ph.D.
 Email: cgaps-remail.nih.gov
 Tissue Procurement: Louis M. Staudt, M.D., Ph.D., David Allman,
 Ph.D., Gerald Marti, M.D.
 cDNA Library Preparation: M. Bento Soares, Ph.D., M. Fatima
 Bonaldo, Ph.D.
 cDNA Library Arrayed by: Greg Lennon, Ph.D.
 DNA Sequencing by: Washington University Genome Sequencing Center
 Clone distribution: NCI-CGAP clone distribution information can be
 found through the I.M.A.G.E. Consortium/LLNL at:
www-bio.llnl.gov/bdip/image/image.html
 Seq primer: -40ml3 fwd. ET from Amersham
 High quality sequence stop: 265.
 FEATURES
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 /mol_type="mRNA"
 /db_xref="taxon:9606"
 /clone="IMAGE:1370071"
 /tissue_type="germinal center B cell"
 /lab_host="DH10B"
 /clone_lib="NCI CGAP GCBI"
 /note="Vector: pT73D-PacI. Site_1: Not I; Site_2: Eco RI;
 1st strand cDNA was prepared from human tonsillar cells
 enriched for germinal center B cells by flow sorting
 (CD20+, IgD-), provided by Dr. Louis M. Staudt (NCI), Dr.
 David Allman (NCI) and Dr. Gerald Marti (CBER). cDNA
 synthesis was primed with a Not I - oligo(dt) primer
 [5'-TGTTACCAATCTGAGTGGAGGCGCGCTCATTTTCTTTT-3'
]. Double-stranded cDNA was ligated to Eco RI adaptor
 (Pharmacia), digested with Not I and cloned into the Not I
 and Eco RI sites of the modified pT73 vector. Library
 went through one round of normalization, and was
 constructed by Bento Soares and M. Fatima Bonaldo."
 ORIGIN
 Query Match 36.4%; Score 145.6; DB 1; Length 323;
 Best Local Similarity 76.0%; Pred. No. 5.5e-18;
 Matches 206; Conservative 1; Mismatches 56; Indels 8; Gaps 2;
 Db 81 ATGCTGTATATCCGACGACTTGGGAGGCCGAGGATCAGTCAAGTCAAGA 140
 Db 284 ATGCTGTATATCCGACGACTTGGGAGGCCGAGGATCAGTCAAGTCAAGT 225
 QY 141 TCGAGACCATCTGGGCAACATGTGAACCCCGTCTTACTTAAATAACAAAATATAGC 200
 Db 224 TCGAAGACAGCTTGCGCAACATGTGAACCTCAATCTTCAATAAAATACAAAATTAGC 165
 QY 201 TGGGATGTGTGACACACCTGTAGTCCAGCTACTCAGAGCCGAGATTTCAGTAGC 255
 Db 164 TGGGATGTGTGATGCTGTGAATCCAGCTACTAGGAGGCTGAGGACGAGATCA 105
 QY 256 --TGAAGTGGAGAGTGAAGCCGAATTCACAGATCAGAGTGAAGTGAAGTGA 312
 Db 104 CTGTGAACCCAGAGGTGAAGGCTTGCAGTGAAGCCGAGATCACTGAGAGACGAATGAGA 45
 QY 313 CKCGCTCAAAAAACAACAAAAA 343
 Db 44 CTCATCTCAAAAAAAGAAAAA 14

```

RESULT 38
BQ267427/c      416 bp      mRNA      linear      EST 15-UTL-2003
LOCUS           1k07f12.x1 Human insulinoma Homo sapiens cDNA clone IMAGE:5780494
DEFINITION      3', mRNA sequence.
ACCESSION       BQ267427
VERSION         BQ267427.1 GI:20492492
KEYWORDS        EST.
SOURCE          Homo sapiens (human)
ORGANISM        Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini;
Homidae; Homo.
REFERENCE       1 (bases 1 to 416)
AUTHORS         Melton, D., Brown, J., Kenty, G., Permutt, A., Lee, C., Kaestner, K.,
                Lemishka, I., Scaer, M., Brestelli, J., Gradwohl, G., Clifton, S.,
                Hillier, L., Marra, M., Page, D., Wylie, T., Martin, J., Blistein, A.,
                Schmitt, A., Weising, B., Ritzer, E., Ronko, I., Bennett, J.,
                Cardenas, M., Gibbons, M., McCann, R., Cole, R., Tsagarashvili, R.,
                Williams, T., Jackson, Y. and Bowers, Y.
                Endocrine Pancreas Consortium
                Unpublished (2000)
                Contact: Douglas Melton, Klaus H. Kaestner, & Hiroshi Inoue
                Endocrine Pancreas Consortium
                Harvard University, Howard Hughes Medical Institute
                Dept of Molecular and Cellular Biology, 7 Divinity Ave, Cambridge,
                MA 02138
                Tel: 617-495-1812
                Fax: 617-495-8557
                Email: dmelton@biohp.harvard.edu
                Library was constructed by Dr. J. Ferrer in vivo mass-excised to
                pBluescript SK- by Dr. H. Inoue DNA sequencing by: Washington
                University Genome Sequencing Center For information on obtaining a
                clone please contact: Dr. Hiroshi Inoue (hinoue@im.wustl.edu)
                Seq primer: -40UP from G1bco
                High quality sequence stop: 355.
FEATURES
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                /mol_type="mRNA"
                /db_xref="taxon:9606"
                /clone="IMAGE:5780494"
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                /lab_host="DH10B (phage-resistant)"
                /clone_lib="Human Insulinoma"
                /note="Organ: pancreas; Vector: pBluescript SK-; Site 1:
                XhoI; Site 2: EcoRI; Constructed with lambda ZapII system
                (Stratagene) by Dr. J. Ferrer, in vivo mass-excised to
                pBluescript SK- by Dr. H. Inoue following the Washington
                University protocol
                (http://genome.wustl.edu/est/lambda_protocol.shtml).
                Please contact Hiroshi Inoue, MD/PhD for further
                information on this library (Metabolism Division, Permutt
                Laboratory, Washington University School of Medicine, Box
                8127, 660 S Euclid Ave, St. Louis, MO 63110). Note: this
                is a Washington University Pancreas EST project library."
ORIGIN
Query Match      36.4%; Score 145.4; DB 3; Length 416;
Best Local Similarity 78.0%; Pred. No. 5.7e-18;
Matches 202; Conservative 1; Mismatches 47; Indels 9; Gaps 2;
OY              81 ATGCTGTATCCAGCACTTCGGAGGCCAAGTGGCGGATCACTGAGGTCAAGAGA 140
DB              250 ACGCTGTATCTCTGACACTTGGAGGCCAAGTGGCGAGATCACTGAGGTCAAGAGT 191
OY              141 TCGAGACCATCTCGGCAACATGGGAACCCCGCTTACTTAATAAATAAATAAATAGC 200
DB              190 TCGAACAACAGCTGACCAATATGATGAACCCCGCTCTCA-AAAAAATACAAAATTTAGC 132
OY              201 TGGGATGATGGACACACCTGTAGTCCAGCTACTACAGAGCCGAGATTGCACTGAGC 260
DB              131 CGGGCTGTGTCAAGCACTGTATCCAGCTACTACAGAGGCAAGAGTTGCAATGGGC 72

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OY              261 TGAATCCAGAGTACGCCGAATATCAAGATCAAGAGTACAGAGTGAACCKCCTCT 320
DB              71 TGAAGTTCG-----GGCCATTGCACTCCAGCCTGGCGAAAAAGATGAGGCTCTGTCT 20
OY              321 CAAACAACAACAACAACA 339
DB              19 CAAAGAAAAAAAAAAAAA 1
RESULT 39
AOS98684        464 bp      DNA      linear      GSS 08-JUN-1999
LOCUS           HS_5335_B2_E09 SP6E RPCT-11 Human Male BAC Library Homo sapiens
DEFINITION      genomic clone Plate=912 Col=18 Row=J, genomic survey sequence.
ACCESSION       AOS98684
VERSION         AOS98684.1 GI:5029896
KEYWORDS        GSS.
SOURCE          Homo sapiens (human)
ORGANISM        Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini;
Homidae; Homo.
REFERENCE       1 (bases 1 to 464)
AUTHORS         Mahairas, G.G., Wallace, J.C., Smith, K., Swartzell, S., Holzman, T.,
                Keller, A., Shaker, R., Furlong, J., Young, J., Zhao, S., Adams, M.D. and
                Hood, L.
                Sequence-tagged connectors: A sequence approach to mapping and
                scanning the human genome
                Proc. Natl. Acad. Sci. U.S.A. 96 (17), 9739-9744 (1999)
                10449764
                Contact: Mahairas GG, Wallace JC, Hood L
                High Throughput Sequencing Center
                University of Washington
                401 Queen Anne Avenue North, Seattle, WA 98109, USA
                Tel: (206) 616-3618
                Fax: (206) 616-3887
                Email: jwallace@u.washington.edu
                Clones are derived from the human BAC library RPCT-11. For BAC
                library availability, please contact Pieter de Jong
                (pieterdejong.med.buffalo.edu). Clones may be purchased from
                BACPAC Resources (http://bacpac.med.buffalo.edu/ordering_bac.htm)
                or from Research Genetics (info@resgen.com). BAC end Web Server:
                http://www.hnsc.washington.edu
                Plate: 912 Row: J Column: 18
                Seq primer: SP6
                Class: BAC ends
                High quality sequence stop: 464.
FEATURES
source          1..464
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                /mol_type="genomic DNA"
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                /clone="Plate=912 Col=18 Row=J"
                /sex="male"
                /clone_lib="RPCT-11 Human Male BAC Library"
                /note="Vector: pBACe3.6; Site 1: EcoRI; Site 2: EcoRI;
                Male blood DNA was isolated from one randomly chosen donor
                and partially digested with a combination of EcoRI and
                EcoRI Methylase. Size selected DNA was cloned into the
                pBACe3.6 vector at EcoRI sites"
ORIGIN
Query Match      36.4%; Score 145.4; DB 11; Length 464;
Best Local Similarity 72.6%; Pred. No. 5.6e-18;
Matches 204; Conservative 1; Mismatches 67; Indels 9; Gaps 1;
OY              81 ATGCTGTATCCAGCACTTCGGAGGCCAAGTGGCGGATCACTGAGGTCAAGAGA 140
DB              97 ATGCTGTATCCAGCACTTGGAGGCCAAGTGGCGGATCACTGAGGTCAAGAGC 156
OY              141 TCGAGACCATCTCGGCAACATGGGAACCCCGCTTACTTAATAAATAAATAAATAGC 200

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Query Match 36.4%; Score 145.4; DB 9; Length 551;
 Best Local Similarity 71.5%; Pred. No. 5.4e-18;
 Matches 191; Conservative 0; Mismatches 76; Indels 0; Gaps 0;

DB 81 ATGCTGTATATCCAGCACTTGGGAGGCCAAGTGGGCGGATCACTTGGTCAAGAGA 140
 DB 392 AGGCTATATATCCAGCACTTGGGAGGCCGCGGTGATCACTTGGTCAAGAGA 333
 DB 141 TCGAATCATCTTGGGCAATGATGAAACCCCTTTACTTAAATAATCAAAAAATAGC 200
 DB 332 TCAACACCAAGCTTGGGCAACACAGTGAACCCCATCTTCAATAAATAATCAAAAAATAGC 273

ORIGIN
 Query Match 36.4%; Score 145.2; DB 10; Length 324;
 Best Local Similarity 71.1%; Pred. No. 6.6e-18;
 Matches 209; Conservative 0; Mismatches 78; Indels 7; Gaps 1;

DB 81 ATGCTGTATATCCAGCACTTGGGAGGCCAAGTGGGCGGATCACTTGGTCAAGAGA 140
 DB 24 ATGCTGTATATCCAGCACTTGGGAGGCCGCGGTGATCACTTGGTCAAGAGA 83
 DB 141 TCGAATCATCTTGGGCAATGATGAAACCCCTTTACTTAAATAATCAAAAAATAGC 200
 DB 84 TCGAATCATCTTGGTCAACATGATGAAACCCCTTCTACTTAAATAATCAAAAAATAGC 143
 DB 201 TGGGATGTGGGACACACTGTATGTCCAGCTACTAGCA-----GCCGAGATATGC 253

RESULT 42
 DM412368 324 bp mRNA linear EST 14-JAN-2006
 LOCUS HHAGE013098 Human liver regeneration after partial hepatectomy Homo
 DEFINITION sapiens cDNA, mRNA sequence.
 ACCESSION DM412368
 VERSION DM412368.1 GI:84913924
 KEYWORDS EST.
 SOURCE Homo sapiens (human)
 ORGANISM Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini;
 Homnidae; Homo.
 REFERENCE 1 (bases 1 to 324)
 Xu, C.S.
 Liver regeneration after PH
 Unpublished (2003)
 JOURNAL Contact: Cun-Shuan Xu
 Henan Bioengineering Key Lab
 Henan Normal University
 No. 148 Jianshe Road, Xinxiang City, P.R.China
 Tel: 0086373328084
 Fax: 0086373326524
 Email: xucs@x263.net..

FEATURES
 source Location/Qualifiers
 1..324
 /organism="Homo sapiens"
 /mol_type="mRNA"
 /db_xref="taxon:9606"
 /tissue_type="liver"
 /clone_lib="Human liver regeneration after partial
 hepatectomy"

DB 144 CGAGGTGTGTGTGACCGCTTATATCCAGTACTTCGGGAGGCTGAGGCAAGAAATTCG 203
 DB 254 AGTGAGCTGAGATCGCAGATGAGGCCGAATCAAGATCAAGATGAGAGTGAAGC 313
 DB 204 CTGTAACCTGGGAGGACAGAGGTTGCAACCATGTGCACTCCAGCTGGCGACAAAGAGCAAG 263

ORIGIN
 Query Match 36.3%; Score 145.2; DB 9; Length 498;
 Best Local Similarity 72.1%; Pred. No. 6.1e-18;
 Matches 189; Conservative 0; Mismatches 73; Indels 0; Gaps 0;

DB 2 CAGGATCATGAGCTGATGCTGGGCGGATGGGAAACCAATATTATTAAGACATTCAGGC 61
 DB 341 CATTTCTCATGATCATTAATGCTCATTTGTATCATCATCTCTTGGAATGCTTAAAAA 282
 DB 62 CAGGATGACATGCTGATGCTGATATCCAGCACTTGGGAGGCCAAGTGGGCGG 121
 DB 281 TGGCCGGGACAGGAGGCTGACAGGCTGTATATCCAGCACTTGGGAGGCCAAGTGGGCGG 222

RESULT 43
 DB330188 498 bp mRNA linear EST 10-DEC-2005
 LOCUS DB330188 PROST2 Homo sapiens cDNA clone PROST2015b1 3', mRNA
 DEFINITION sequence.
 ACCESSION DB330188
 VERSION DB330188.1 GI:83456206
 KEYWORDS EST.
 SOURCE Homo sapiens (human)
 ORGANISM Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini;
 Homnidae; Homo.
 REFERENCE 1 (bases 1 to 498)
 Kimura, K., Wakamatsu, A., Suzuki, Y., Ota, T., Nishikawa, T.,
 Yamashita, R., Yamamoto, J., Sekine, M., Teuriltan, K., Wakaguri, H.,
 Ishii, S., Sugiyama, T., Saito, K., Isono, Y., Irie, R., Kushida, N.,
 Yoneyama, T., Otsuka, R., Kanda, K., Yokoi, T., Kondo, H., Wagatsuma, M.,
 Murakawa, K., Ishida, S., Iehibashi, T., Takahashi-Fujii, A., Sugano, S.,
 Tanase, T., Nagai, K., Kikuchi, H., Nakai, K., Isegai, T. and Sugano, S.
 Diversification of Transcriptional Modulation: Large-scale
 Identification and Characterization of Putative Alternative
 Promoters of Human Genes
 JOURNAL Genome Res. 16 (1), 55-65 (2006)
 PUBMED 16344560
 COMMENT Contact: Takao Isegai
 Fij Project (HRI Team)
 Helix Research Institute
 2-6-7 Kazusa-Kamatari, Kisarazu, Chiba, 292-0818, Japan
 Tel: 81-438-52-3975
 Fax: 81-438-52-3986
 Email: fji@cnaeaffity.com
 NEDO human cDNA project (New Energy and Industrial Technology
 Developmental Organization, Japan); cDNA library construction;
 Helix Research Institute (HRI); 5'-end one pass sequencing: HRI,
 Research Association for Biotechnology (RAB) and Biotechnology
 Center, National Institute of Technology and Evaluation; 3'-end one
 pass sequencing: RAB.
 FEATURES
 source Location/Qualifiers
 1..498
 /organism="Homo sapiens"
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 /db_xref="taxon:9606"
 /clone="PROST2015b1"
 /tissue_type="prostate"
 /clone_lib="PROST2"
 /note="Vector: pME18SFL3"

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QY 122 ATGACCTGAGGTGACAGATGAGACCATCTCTGGCCAAACATGTGAAACCCCGTCTTAC 181
DB 221 ATGACCTGAGGTGAGGAGTTGAGACGATCTGGCCAAATGAGAAACCCCGTCTTAC 162
QY 182 TAAATAATACAAAATAGTCTGGGATGTGGACACACCTGTACTCCAGCTACTCAGGA 241
DB 161 TAAATAATACAAAATAGTCTGGGATGTGGGATGTGGGATGTGGGATGTGGGATGTGGG 102
QY 242 GCCGAGATTTGCACTGAGCTGA 263
DB 101 GTCAGAGACTGGGTTCAATTGA 80

RESULT 44
DA714821/c 515 bp mRNA linear EST 13-NOV-2005
LOCUS DA714821 NT2R12 Homo sapiens cDNA clone NT2R12021923 5', mRNA
DEFINITION sequence.
ACCESSION DA714821
VERSION DA714821.1 GI:82370722
KEYWORDS EST.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini;
Hominiidae; Homo.
REFERENCE 1 (bases 1 to 515)
AUTHORS Kimura,K., Wakamatsu,A., Suzuki,Y., Ota,T., Nishikawa,T.,
Yamashita,R., Yamamoto,J., Sekine,M., Tsuritani,K., Wakaguri,H.,
Ishii,S., Sugiyama,T., Saito,K., Isono,Y., Irie,R., Kushida,N.,
Yonekawa,T., Otsuka,R., Kanda,K., Yokoi,T., Kondo,H., Matsushima,M.,
Morakawa,K., Ishida,S., Ishibashi,T., Takahashi-Fujii,A.,
Tanase,T., Nagai,K., Kikuchi,H., Nakai,K., Isegaki,T. and Sugano,S.
Diversification of Transcriptional Modulation: Large-scale
Identification and Characterization of Putative Alternative
Promoters of Human Genes
JOURNAL Genome Res. 16 (1), 55-65 (2006)
PUBMED 16344560
COMMENT Contact: Takao Isegaki
FUJ Project (HRI Team)
Helix Research Institute
2-6-7 Kazusa-Kamatari, Kisarazu, Chiba, 292-0818, Japan
Tel: 81-438-52-3975
Fax: 81-438-52-3986
Email: fuj-cdna@hri.fcy.com
NEO human cDNA project (New Energy and Industrial Technology
Developmental Organization, Japan); cDNA library construction:
Helix Research Institute (HRI); 5'-end one pass sequencing: HRI,
Research Association for Biotechnology (RAB) and Biotechnology
Center, National Institute of Technology and Evaluation; 3'-end one
pass sequencing: RAB.
Location/Qualifiers
1..515
/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/clone="NT2R12021923"
/cell_line="tetatocarcinoma"
/clone_lib="NT2"
/clone_1ib="NT2R12"
/clone_2ib="PME18SFL3"
/feature="vector: pME18SFL3; majorly NT2 neuron; mRNA from
NT2 neuronal precursor cells treated 2-weeks mitotic
inhibitor after 5-weeks retinoic acid (RA) induction."

ORIGIN
Query Match 36.3%; Score 145.2; DB 9; Length 515;
Best Local Similarity 78.4%; Pred. No. 6e-18; Indels 0; Gaps 0;
Matches 174; Conservative 0; Mismatches 48; Indels 0; Gaps 0;

QY 81 ATGCTGTAATCCAGCACTTCGGAGGCGAAGTGGGCGATCACTGAGGTCAAGGA 140
DB 226 ACGCTATATATCCAGCACTTGGAGGCGTGGGCGGATGATCACTGAGGTCAAGAGT 167

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QY 141 TCGAGACCATCTCGGCCAATGATGTAACCCCGTCTTTCTAATAAATAAATAATAGC 200
DB 166 TCAAGACCACTCTGGCCAAACAGTGAACCCCATCTCTAATAAATAAATAATAGC 107
QY 201 TGGGCATGTGGGACACACCTGTATGCCAGTACTCAGAGCCCGAGATTGCACTGAC 260
DB 106 TGGGCATGATGTGGGACACCTGTATGCCAGTACTCAGAGCCCGAGATTGCACTGAC 47
QY 261 TGAATCGCAGAGTGAACCCGAATCAAGATCAAGATGAG 302
DB 46 CAAGATCGACAACTGCATTCGACCTGGGTGAGAGAGCAAG 5

RESULT 45
AQ314853/c 555 bp DNA linear GSS 04-MAY-1999
LOCUS AQ314853
DEFINITION R0111-96D8-TV R0111-11 Homo sapiens genomic clone R0111-96D8,
genomic survey sequence.
ACCESSION AQ314853
VERSION AQ314853.1 GI:4046316
KEYWORDS GSS.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini;
Hominiidae; Homo.
REFERENCE 1 (bases 1 to 555)
AUTHORS Adams,M.D., Rounsley,S.D., Zhao,S., Baas,S., Linher,K., Golden,K.,
Berry,K., Granger,D., Suh,E., Wible,C., de Jong,F. and Venter,J.C.
Use of human BAC End Sequences for Sequence-Ready Map Building
Unpublished (1998)
JOURNAL Other GSSs: R0111-96D8-TV
COMMENT Contact: Shaying Zhao, William Nierman, Mark Adams
Department of Eukaryotic Genomics
The Institute for Genomic Research
9712 Medical Center Dr., Rockville, MD 20850
Tel: 301 838 0200
Fax: 301 838 0208
Email: hbe@tigr.org
Clones are derived from the human BAC library R0111-11. For BAC
library availability, please contact Pieter de Jong
(pieter@dejong.med.buffalo.edu). Clones may be purchased from
BACPAC Resources (http://bacpac.med.buffalo.edu/ordering) or from
Research Genetics (info@resgen.com). BAC end search page:
http://www.tigr.org/tcd/humgen/bac_end_search/bac_end_search.html
Seq primer: T7
Class: BAC ends.
Location/Qualifiers
1..555
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/mol_type="genomic DNA"
/db_xref="GDB:7536559"
/db_xref="taxon:9606"
/clone="R0111-96D8"
/sex="Male"
/cell_type="Lymphocytes"
/clone_lib="R0111"
/feature="vector: pBAC3.6; Site_1: EcoRI; Site_2: EcoRI;
R0111 Human Male BAC Library"

ORIGIN
Query Match 36.3%; Score 145.2; DB 11; Length 555;
Best Local Similarity 73.4%; Pred. No. 5.9e-18; Indels 14; Gaps 1;
Matches 204; Conservative 1; Mismatches 59; Indels 14; Gaps 1;

QY 84 CCTGTAATCCAGCACTTCGGAGGCGAAGTGGGCGATCACTGAGGTCAAGATCG 143
DB 374 CCTGTAATCCAGCACTTCGGAGGCGAAGTGGGCGATCACTGAGGTCAAGATCG 315
QY 144 AGACCATCTCGGCCAATGATGTAACCCCGTCTTTCTAATAAATAAATAATAGCTG 203
DB 314 AGACCATCTCGGCCAATGATGTAACCCCGTCTTTCTAATAAATAAATAATAGCTG 255

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QY 204 GCATGTGGACACACTTGTAGTCCAGTACTACAGAGCCGAGATTGTCAGTACGTGA 263
 DB 254 GCGTGGTGAACGACCTTGTAGTCCAGTACTCCGAGGCTGAGGACAGAGAAATGCTT 195
 QY 264 GATCGCAG-----AGTAGCCGAAATCACAGATCACAGAGTGAAGCAGAGTG 309
 DB 194 GAACCCAGAGGCGGAGTTGCAGTGAGCCGAGATCACGCCACCTGATTCAGTCCAGTG 135
 QY 310 AGACGCCGCTCAAAAACAACAACAAAAA 347
 DB 134 AGACTCGCTCAAAAAAAGTAAGTTAATTA 97
 RESULT 46
 CB309712 591 bp mRNA linear EST 26-MAR-2004
 LOCUS CB309712
 DEFINITION AGENCOURT_11828537 NICHD_Rh_Ov1 Macaca mulatta cDNA clone
 IMAGE:6913528 5', mRNA sequence.
 ACCESSION CB309712
 VERSION CB309712
 KEYWORDS CB309712.1 GI:28832422
 SOURCE EST
 ORGANISM Macaca mulatta (rhesus monkey)
 Macaca mulatta
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini;
 Cercopithecoidea; Cercopithecinae; Macaca.
 1 (bases 1 to 591)
 NCI-CCAP http://www.ncbi.nlm.nih.gov/ncicgap.
 National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
 Tumor Gene Index
 Unpublished (1997)
 JOURNAL Contact: Daniela S. Gerhard, Ph.D.
 COMMENT Office of Cancer Genomics
 National Cancer Institute / NIH
 Bldg. 31 Rm10A07 Bethesda, MD 20892
 Email: cgapbs-remail.nih.gov
 Tissue Procurement: Dr. Eliot Spindel
 CDNA Library Preparation: CLONTECH
 DNA Sequencing by: The I.M.A.G.E. Consortium (ILMIL)
 Clone distribution: NCI-CGAP clone distribution information can be
 found through the I.M.A.G.E. Consortium/ILMIL at:
 http://image.llnl.gov
 Plate: L10CM3154 row: m column: 15
 High quality sequence stop: 454.
 Location/Qualifiers
 1..591
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 /clone="IMAGE:6913528"
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 /lab_host="DH10B (phage-resistant)"
 /clone_lib="NICHD Rh_Ov1"
 /note="Organ: ovary; Vector: pDNR-LIB; Site 1: Sfi I;
 Site 2: Sfi I; cloned unidirectionally. Primer: Oligo dT.
 Average insert size 1.0-4.0 Kb. Tissue pooled from
 pre-puberatal, post pubertal sn menopausal monkeys.
 Constructed by Clontech. Note: this is a NICHD Library."
 ORIGIN
 Query Match 36.3%; Score 145.2; DB 4; Length 591;
 Best Local Similarity 84.8%; Pred. No. 5.9e-18;
 Matches 162; Conservative 0; Mismatches 29; Indels 0; Gaps 0;
 QY 81 ATGCTGTATATCCAGCACTTCGAGAGCCCAAGTGGCGATCCTGAGGTCAAGAGA 140
 DB 287 ATGCTGTATATCCAGCACTTCGAGAGCCCAAGTGGCGATCCTGAGGTCAAGAGA 346
 QY 141 TCGAGACATCTCGGCGCAATGCTGAATCCCGCTTTACTAAATAATCAAAAAATAGC 200
 DB 347 TCAAGACCAACCTGGCGCAATGCTGAATCCCGCTTTACTAAATAATCAAAAAATAGC 406

QY 201 TGGGATGTGGACACACTTGTAGTCCAGTACTACAGAGCCGAGATTGTCAGTACG 260
 DB 407 TGGGATGTGGACACACTTGTATCCAGTACTCAGAGGCTGAGGACAGAGATCC 466
 QY 261 TGAGATCGCAG 271
 DB 467 CTGGAACCCAG 477
 RESULT 47
 DA408951 649 bp mRNA linear EST 07-NOV-2005
 LOCUS DA408951
 DEFINITION DA408951 BRTHA3 Homo sapiens cDNA clone BRTHA3015134 5', mRNA
 sequence.
 ACCESSION DA408951
 VERSION DA408951
 KEYWORDS DA408951.1 GI:81121486
 SOURCE EST
 ORGANISM Homo sapiens (human)
 Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini;
 Homnidae; Homo.
 1 (bases 1 to 649)
 Kimura,K., Wakamatsu,A., Suzuki,Y., Ota,T., Nishikawa,T.,
 Yamashita,R., Yamamoto,J., Sekine,M., Tsurikani,K., Wakaguri,H.,
 Ishii,S., Sugiyama,T., Saito,K., Isono,Y., Irie,R., Kushida,N.,
 Yoneyama,T., Otsuka,R., Kanda,K., Yokoi,T., Kondo,H., Wagatsuma,M.,
 Murakawa,K., Ishida,S., Ishibashi,T., Takahashi-Fujii,A.,
 Tanase,T., Nagai,K., Kikuchi,H., Nakai,K., Isogai,T. and Sugano,S.
 Diversification of Transcriptional Modulation: Large-scale
 Identification and Characterization of Putative Alternative
 Promoters of Human Genes
 JOURNAL Genome Res. 16 (1), 55-65 (2006)
 COMMENT 16344560
 PUBLISHED
 CONTACT: Takao Isogai
 FJI Project (HRI Team)
 Helix Research Institute
 2-6-7 Kazusa-Kamatari, Kisarazu, Chiba, 292-0818, Japan
 Tel: 81-438-52-3975
 Fax: 81-438-52-3986
 Email: fji-cdna@nifty.com
 NEDO human cDNA project (New Energy and Industrial Technology
 Developmental Organization, Japan); cDNA library construction:
 Helix Research Institute (HRI); 5'-end one pass sequencing: HRI,
 Research Association for Biotechnology (RAB) and Biotechnology
 Center, National Institute of Technology and Evaluation; 3'-end one
 pass sequencing: RAB.
 Location/Qualifiers
 1..649
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 /db_xref="taxon:9606"
 /clone="BRTHA3015134"
 /tissue_type="thalamus"
 /clone_lib="BRTHA3"
 /note="Vector: pME18SFL3"
 ORIGIN
 Query Match 36.3%; Score 145.2; DB 9; Length 649;
 Best Local Similarity 70.1%; Pred. No. 5.8e-18;
 Matches 195; Conservative 0; Mismatches 83; Indels 0; Gaps 0;
 QY 26 CATGGAAACCAATATTAATTAAGACATTTGTCAGGCCGAGGATGACACTGCTGAATGC 85
 DB 273 CCGGACCACTCCCTCTTAATTAAGAAATAGAAATAGGCGCAAGCAGATGCTCATGTC 332
 QY 86 TGTATCCCGCACTTGGGAGCGCAAGTGGCGGATCCTGAGGTCAAGAGTCAAGATGAG 145
 DB 333 TGTATCTCAGCACTTGGGAGCGCTGAGGTGAGAGATCACTTAAGGTCAAGATTTCCAG 392
 QY 146 ACCATCTGGCGCAATGCTGAATCCCGCTTTACTAAATAATCAAAAAATAGTGGGC 205
 DB 393 ACCAGCCGCGCAATGCTGAGCAATCCCGCTCTTAATAATAATCAAAAAATAGTGGGC 452

QY	206	ATGTTGGGACACACCTGTAGTCCAGGACACCTGAGAGCCGAGATGTGAGTGAAGACTGAGA	265
Db	453	GTGTTGTAACACACCTGTATTCACAGGCTACTGGGAGGCTTAGGACGAGATTGCTTGA	512
QY	266	TCGACAGATGAGCCGAATTCACAGATCAAGAGTGAGC	303
Db	513	GCCAGAAAGGCGAGGTTGCAGTGAACCGAATTATGC	550
RESULT 48			
LOCUS	C2459725	849 bp	DNA
DEFINITION	MCF748c16TF Human MCF7 breast cancer cell line library (MCF7_1)		linear GSS 20-OCT-2005
ACCESSION	C2459725		
VERSION	C2459725.1		
KEYWORDS	GI:77937953		
SOURCE	GSS.		
ORGANISM	Homo sapiens (human)		
	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini; Homiidae; Homo.		
REFERENCE	1 (bases 1 to 849)		
AUTHORS	Volik,S.V., Raphael,B.J., Huang,G.-Q., Murnane,J., Brehner,J.H., Bajsarowicz,K., Paris,P., Tao,Q., Kowbel,D., Lapuk,A.V., Kuo,W.-L., Shagin,D.A., Shagina,I.A., Magrane,G., Gray,J.W., Jan,F.-C., de Jong,P., Pevzner,P. and Collins,C.		
TITLE	Decoding the genomic architecture and high throughput detection of fusion transcripts in breast cancer cell lines: implications for a tumor genome project		
JOURNAL	Unpublished (2005)		
COMMENT	Contact: Volik SV Colin Collins' lab UCSF Comprehensive Cancer Center UCSF Box 0808, San Francisco, CA 94143-0808, USA Tel: 415 502 7066 Fax: 415 502 5665 Email: svolik@cc.ucsf.edu This clone is available from Amplicon Express http://www.genomex.com Class: BAC ends.		
FEATURES			
source	Location/Qualifiers 1..849 /organism="Homo sapiens" /mol_type="genomic DNA" /db_xref="taxon:9606" /clone="MCF7_48c16" /sex="Female" /clone_11b="Human MCF7 breast cancer cell line library (MCF7_1)" /note="Vector: pECBAC1; Site 1: HindIII; This library was constructed from MCF7 breast cancer cell line by Amplicon Express (http://www.genomex.com) using their standard procedure."		
ORIGIN			
Query Match	36.3%	Score 145.2	DB 13; Length 849;
Best Local Similarity	73.4%	Pred. No. 5.5e-18;	
Matches 204; Conservative	1; Mismatches 59; Indels 14; Gaps 1;		
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ORGANISM	Homo sapiens		
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AUTHORS	1 (bases 1 to 550) Kamura,K., Wakamatsu,A., Suzuki,Y., Ota,T., Nishikawa,T., Yamashita,R., Yamamoto,J., Sekine,M., Tsutitani,K., Wakaguri,H., Ishii,S., Sugiyama,T., Saito,K., Isono,Y., Irie,R., Kushida,N., Yoneyama,T., Otsuka,R., Kaneda,K., Yokoi,T., Kondo,H., Magatsuna,M., Murakawa,K., Ishida,S., Iihashi,T., Takahashi-Fujii,A., Tanase,T., Nagai,K., Kikuchi,H., Nakai,K., Isogai,T. and Sugano,S.		
TITLE	Diversification of Transcriptional Modulation: Large-scale Identification and Characterization of Putative Alternative Promoters of Human Genes		
JOURNAL	Genome Res. 16 (1), 55-65 (2006)		
PUBMED	16344560		
COMMENT	Contact: Takao Isogai FLU Project (HRI Team) Helix Research Institute 2-6-7 Kazusa-Kamatari, Kisarazu, Chiba, 292-0818, Japan Tel.: 81-438-52-3975 Fax: 81-438-52-3986 Email: flj@cnaomifly.com NEO human cDNA project (New Energy and Industrial Technology Developmental Organization, Japan); cDNA library construction: Helix Research Institute (HRI); 5'-end one pass sequencing: HRI, Research Association for Biotechnology (RAB) and Biotechnology Center, National Institute of Technology and Evaluation; 3'-end one pass sequencing: RAB. Location/Qualifiers 1..550 /organism="Homo sapiens" /mol_type="mRNA" /db_xref="taxon:9606" /clone="BRACE3J030866" /tissue type="cerebellum" /clone_id="BRACE3" /note="Vector: pME18SFL3"		
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DEFINITION DA198015 BRAH2 Homo sapiens cDNA clone BRAH2000862 5', mRNA
 LOCUS sequence.

ACCESSION DA198015

VERSION DA198015.1 GI:78309617

KEYWORDS EST.

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 ORGANISM Homo sapiens
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 Homnidae; Homo.

REFERENCE 1 (bases 1 to 555)
 Kimura,K., Wakamatsu,A., Suzuki,Y., Ota,T., Nishikawa,T.,
 Yamashita,R., Yamamoto,J., Sekine,M., Tsuritani,K., Wakaguri,H.,
 Ichii,S., Sugiyama,T., Saito,K., Isomoto,Y., Irie,R., Kishida,N.,
 Yoneyama,T., Otsuka,K., Kanda,K., Yokoi,T., Kondo,H., Wagatsuma,M.,
 Murakawa,K., Ishida,S., Ishibashi,T., Takahashi-Fujii,A.,
 Tanase,T., Nagai,K., Kikuchi,H., Nakai,K., Isogai,T. and Sugano,S.
 Diversification and Transcriptional Modulation: Large-scale
 Identification and Characterization of Putative Alternative
 Promoters of Human Genes
 Genome Res. 16 (1), 55-65 (2006)

TITLE

JOURNAL PUBMED
 16344560

COMMENT

Contact: Takao Isogai
 Fij Project (HRI Team)
 Helix Research Institute
 2-6-7 Kazusa-Kamatari, Kisarazu, Chiba, 292-0818, Japan
 Tel: 81-438-52-3975
 Fax: 81-438-52-3986
 Email: filj-cdna@nifty.com
 NEDO human cDNA project (New Energy and Industrial Technology
 Developmental Organization, Japan); cDNA library construction:
 Helix Research Institute (HRI); 5'-end one pass sequencing: HRI,
 Research Association for Biotechnology (RAB) and Biotechnology
 Center, National Institute of Technology and Evaluation; 3'-end one
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DB 367 CCGCATGTGTGCGCACACCTGTAAATCCAGCTACTCGAAGCTGTGAGCAGAGAG 311

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 Job time : 4693 secs

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GenCore version 5.1.9
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On nucleic - nucleic search, using sw model

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Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 150 summaries

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Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

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ALIGNMENTS

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; Patent No. 6812339
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; APPLICANT: VENTER, J. Craig et al.
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; WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
; FILE REFERENCE: C1001307
; CURRENT APPLICATION NUMBER: US/09/949,016
; CURRENT FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
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; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FASTSEQ for Windows Version 4.0
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; ORGANISM: Human
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141 TCGAGACATCTGCGCAACATGAGTGAACCCCGCTTTACTATAAATACAAATATAGC 200
505 TCAAGATCAGCTTCCGCAACATGAGTGAACCCCGCTTTACTATAAATATAGC 446
201 TGGCATGTGCGCACACACTGTAGTCCAGCTTCTCAGAGCCGAGATTGCAATGAC 260
445 TGGCATGTGCGCGGCGGAGCTGAGTCCAGCTTCTCAGAGCGGAGGAGCAATGCG 386
261 TGAATGCGCAG-----AGTAGCCGAATTCACATCAGATGAGCAGA 306
385 CTTAACCCAGCGCGGAGGTTGACAGTCCAGATGACACAGACTTGGTATCAGAC 326
307 GTGAGACGCGCTCAAAAACACACAAACAAACAAACAAACAAACAAACAAAC 354
325 GTGAGACTTCACTCAAAAACAAACAAACAAACAAACAAACAAACAAACAAAC 278
```

```
RESULT 2
US-09-949-016-14471/c
; Sequence 14471, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
; FILE REFERENCE: C1001307
; CURRENT APPLICATION NUMBER: US/09/949,016
; CURRENT FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FASTSEQ for Windows Version 4.0
; SEQ ID NO 14471
; LENGTH: 10980
; TYPE: DNA
; ORGANISM: Human
US-09-949-016-14471
```

```
Query Match 39.6%; Score 158.4; DB 3; Length 10980;
Best Local Similarity 73.9%; Pred. No. 9e-37;
Matches 201; Conservative 0; Mismatches 71; Indels 0; Gaps 0;
```

```
76 GCTGAATCCGTGTATCCAGACACTTCCGAGCGCAAGTGGCGGATCACTGAGTCA 135
4821 GATTACCGCTGTATCCAGACACTTCCGAGCGCGAGCGGAGTCACTGAGTCA 4762
136 AGAATCGACACATCCGCGCAACATGAGTGAACCCCGCTTTACTATAAATACAAA 195
4761 GAGATTGAGTCACTGCGCAACATGAGTGAACCCCGCTTTACTATAAATACAAA 4702
```

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QY 196 ATAGCTGGGATGTGGGACACACCTGTATCCAGTACTGACGAGCGGAGATTGAC 255
DB 4701 CCAAGCTGGGTGGTGGTACACCTTTGATCCAGCTACTCTGGAGCGGAGTTGAC 4642
QY 256 TGAAGTGAATGACGAGAGTACCGAATACAGATCAGAGTGAAGTGAAGAC 315
DB 4641 TGAAGCGAAATGACCACTACCTCCAGCTGGGGTGAAGAGGAGACTGTCTCCCA 4582
QY 316 CGTCTCAAAAACACACACAAAAA 347
DB 4581 GGAAGAAAAA 4550

```

RESULT 3

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US-09-949-016-12783/c
; Sequence 12783, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
; FILE REFERENCE: CL001307
; CURRENT APPLICATION NUMBER: US/09/949,016
; PRIOR FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 12783
; LENGTH: 15564
; TYPE: DNA
; ORGANISM: Human
US-09-949-016-12783

```

```

Query Match 39.6%; Score 158.4; DB 3; Length 15564;
Best Local Similarity 73.9%; Pred. No. 1e-36;
Matches 201; Conservative 0; Mismatches 71; Indels 0; Gaps 0;

```

```

QY 76 GGTGATGCTGTATCCAGACACTTCGGAGGCGCAAGTGGCGGAGTCACTGAGTCA 135
DB 9414 GGTGATGCTGTATCCAGACACTTCGGAGGCGGAGTGGAGTCACTGAGTCA 9355
QY 136 AGAGATCGAGACCTCTGGCCACATGTGTGAACCCCGCTTTACTAATAACAAAA 195
DB 9354 GGAAGTTGATGATCAGTCTGGCCACATGTGTGAACCCCGCTTTACTAATAACAAAA 9295
QY 196 ATAGCTGGGATGTGGGACACACCTGTATCCAGTACTGACGAGCGGAGATTGAC 255
DB 9294 CCAAGCTGGGTGGTGGTACACGCTTTGATCCAGCTACTCTGGAGGCGGAGTTGAC 9235
QY 256 TGAAGTGAATGACGAGAGTACCGAATACAGATCAGAGTGAAGTGAAGAC 315
DB 9234 TGAAGCGAAATGACCACTACCTCCAGCTGGGGTGAAGAGGAGACTGTCTCCCA 9175
QY 316 CGTCTCAAAAACACACAAAAA 347
DB 9174 GGAAGAAAAA 4550

```

RESULT 4

```

US-09-949-016-17230
; Sequence 17230, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
; FILE REFERENCE: CL001307
; CURRENT APPLICATION NUMBER: US/09/949,016

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; CURRENT FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 17230
; LENGTH: 90776
; TYPE: DNA
; ORGANISM: Human
US-09-949-016-17230

```

```

Query Match 39.6%; Score 158.4; DB 3; Length 90776;
Best Local Similarity 75.0%; Pred. No. 2.1e-36;
Matches 216; Conservative 1; Mismatches 57; Indels 14; Gaps 1;

```

```

QY 81 ATGCTGTATATCCAGACCTTCGGAGGCGCAAGTGGCGGAGTCACTGAGTCAAGAA 140
DB 10217 ACGCTGTATATCCAGACCTTCGGAGGCGGAGGCGGAGTTCCTGAGGTCAAGAT 10276
QY 141 TGAAGACCATCTGCGCAACATGTGTAACCCCGCTTTACTAATAAATAATAGC 200
DB 10277 TGAAGATAGCTGCGCAACATGTGTAACCCCGCTTTACTAATAAATAATAGC 10336
QY 201 TGGGATGTGGGACACACCTGTATCCAGTACTGACGAGCGGAGATTGACGAGC 260
DB 10337 TGGGATGTGGGCGGACCTGTATCCAGTACTGCGGAGGTGAAGGAGGAGATGC 10396
QY 261 TGAATCCGAC-----AGTGAAGCGGAATACAGATCAGAGTGAAGCAGA 306
DB 10397 CTGGAACCGGCGGAGGAGTTCAGTGAAGCGGAGTTCAGCAGCTTGGGTGCAAGC 10456
QY 307 GTGAGACGCGCTCAAAAACACACAAAAAACCATTAG 354
DB 10457 GTGAGACTTCACTCAAAAACACACAAAAAACACCAAGATG 10504

```

```

RESULT 5
US-09-949-016-13909
; Sequence 13909, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
; FILE REFERENCE: CL001307
; CURRENT APPLICATION NUMBER: US/09/949,016
; CURRENT FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 13909
; LENGTH: 44789
; TYPE: DNA
; ORGANISM: Human
US-09-949-016-13909

```

```

Query Match 38.6%; Score 154.4; DB 3; Length 44789;
Best Local Similarity 72.5%; Pred. No. 2.4e-35;
Matches 200; Conservative 0; Mismatches 76; Indels 0; Gaps 0;

```

```

QY 81 ATGCTGTATATCCAGACCTTCGGAGGCGCAAGTGGCGGAGTCACTGAGTCAAGAA 140
DB 14253 ATGCTGTATATCCAGACCTTCGGAGGCGCAAGTGGGTGATCATTTGGGTGAGAGT 14312

```

```
QY      141  TCGAGACCATCTGGCCCAACATGTTGAACCCCGTCTTACTTAAATAACAAATAATGAC 200
      |||||
Db      1413  TTGAGACCATGCTGGCCCAACATGTTGAACCCCGCTTATTAATAATAACAAATAATGAC 14372
QY      201  TGGGCATGTGTGGCACAACACTGTAGTCCAGCTACTCAAGAGCCGGAGATTGCAGTGAGC 260
      |||||
Db      14373  TGGGCATGTGTGGCACAAGTGTGCTGTAGTCCAGCTACTCAAGAGCCGGAGATTGCAGTGAGT 14432
QY      261  TGAATGTCAGAGTGAAGCCGAATTCACAGATTCACAGATGAGCAGAGTGAACKCCGCT 320
      |||||
Db      14492  TGAATGTGTGCTGCTGCACTGCACTCCAGCTTCGCTGACAGATGAGATGCTGTCTCAAAAAAAA 14492
QY      321  CAAAAACAACAACAAAAAACCATAAGC 356
      |||||
Db      14493  AAAAAACAACAACAAAAAACCATAAGC 14528
```

```
RESULT 6
US-09-949-016-12542/c
; Sequence 12542, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; FILE REFERENCE: C1001307
; CURRENT APPLICATION NUMBER: US/09/949,016
; CURRENT FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 12542
; LENGTH: 173787
; TYPE: DNA
; ORGANISM: Human
US-09-949-016-12542
```

```
Query Match      38.6%; Score 154.4; DB 3; Length 173787;
Best Local Similarity 76.9%; Pred. No. 4.2e-35;
Matches 203; Conservative 1; Mismatches 52; Indels 8; Gaps 1;

QY      81  ATGCTGTAAATCCAGCACTTCGGAGGCCAAGGTGGCCGGATCACTGAGGTCAAGAGA 140
      |||||
Db      18373  ACGCTGTAAATCCAGCGCTTTGGAGGCGCAAGGTGGAGGATCACTGAGGTCAAGAGT 18314
QY      141  TCGAGACCATCTGGCCCAACATGTTGAACCCCGTCTTACTTAAATAACAAATAATGAC 200
      |||||
Db      18313  TTGAGACCATGCTGGCCCAACATGTTGAACCCCGTCTTACCAAAAAATACAAATAATGAC 18254
QY      201  TGGGCATGTGTGGCACAACACTGTAGTCCAGCTACTCAAGAGCCGGAGATTGCAGTGAGC 260
      |||||
Db      18253  TGGGCATGTGTGGCACAACCTGTATCCAGCTACTCAAGAGCCGGAGATTGCAGTGAGC 18194
QY      261  TGAATGTCAGAGTGAAGCCGAATTCACAGATTCACAGATGAGCAGAGTGAACKCCGCT 320
      |||||
Db      18193  TGAGATC-----ATGCCACTGCACTCCAGCTCGGGCGACAGAGCAAAATCTCCACT 18142
QY      321  CAAAAACAACAACAAAAAACCATAAGC 344
      |||||
Db      18141  CAAAGAAAAAAGCCAA 18118
```

```
RESULT 7
US-09-949-016-17302/c
; Sequence 17302, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
```

```
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; TITLE OF INVENTION: WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
; FILE REFERENCE: C1001307
; CURRENT APPLICATION NUMBER: US/09/949,016
; CURRENT FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 17302
; LENGTH: 173791
; TYPE: DNA
; ORGANISM: Human
US-09-949-016-17302
```

```
Query Match      38.6%; Score 154.4; DB 3; Length 173791;
Best Local Similarity 76.9%; Pred. No. 4.2e-35;
Matches 203; Conservative 1; Mismatches 52; Indels 8; Gaps 1;

QY      81  ATGCTGTAAATCCAGCACTTCGGAGGCCAAGGTGGCCGGATCACTGAGGTCAAGAGA 140
      |||||
Db      18373  ACGCTGTAAATCCAGCGCTTTGGAGGCGCAAGGTGGAGGATCACTGAGGTCAAGAGT 18314
QY      141  TCGAGACCATCTGGCCCAACATGTTGAACCCCGTCTTACTTAAATAACAAATAATGAC 200
      |||||
Db      18313  TTGAGACCATGCTGGCCCAACATGTTGAACCCCGTCTTACCAAAAAATACAAATAATGAC 18254
QY      201  TGGGCATGTGTGGCACAACACTGTAGTCCAGCTACTCAAGAGCCGGAGATTGCAGTGAGC 260
      |||||
Db      18253  TGGGCATGTGTGGCACAACCTGTATCCAGCTACTCAAGAGCCGGAGATTGCAGTGAGC 18194
QY      261  TGAATGTCAGAGTGAAGCCGAATTCACAGATTCACAGATGAGCAGAGTGAACKCCGCT 320
      |||||
Db      18193  TGAGATC-----ATGCCACTGCACTCCAGCTCGGGCGACAGAGCAAAATCTCCACT 18142
QY      321  CAAAAACAACAACAAAAAACCATAAGC 344
      |||||
Db      18141  CAAAGAAAAAAGCCAA 18118
```

```
RESULT 8
US-09-949-016-15580/c
; Sequence 15580, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; FILE REFERENCE: C1001307
; CURRENT APPLICATION NUMBER: US/09/949,016
; CURRENT FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 15580
; LENGTH: 71574
; TYPE: DNA
; ORGANISM: Human
; FEATURES:
; NAME/KEY: misc feature
; LOCATION: (1)...(71574)
; OTHER INFORMATION: n = A,T,C or G
US-09-949-016-15580
```

Query Match 38.4%; Score 153.6; DB 3; Length 71574;
Best Local Similarity 72.8%; Pred. No. 5.1e-35;
Matches 198; Conservative 0; Mismatches 74; Indels 0; Gaps 0;

QY 81 ATGCTGTAATCCAGCACTTGGGAGGCCAAGTGGCGGATCACTGAGTCAAGAGA 140
DB 57890 ACGCTGTAATCTCACTTTGGGAGGCTGAAGTGGCGGATCACTGAGTCAAGAGA 140
QY 141 TCGAGACCACTCGGCGCAACATGATGAAACCCCGCTTTACTTAAATAATCAAAAAATGCG 200
DB 57830 TCGAACCAAGCTGGCGCAACATGATGAAACCCCGCTTTACTTAAATAATCAAAAAATGAGC 57771
QY 201 TGGGATGATGCGACACACTGTAGTCCAGCTACTCAGAGACCGGAGATTGAGTGAAGC 260
DB 57770 TGGGATGATGATGCGGCGCTGTATCCAGCTACTCAGAGACCGGAGATTGAGTGAAGC 57711
QY 261 TGAGATCGCAGATGAGCCGAATATCAAGATCAAGAGTGAAGCAGAGTGAAGCAGCCTGCT 320
DB 57710 TGAGATGATGCGCAGCTGCACTCCAGCTGGGTGACAGAGAACCTGCTCAAAAAA 57651
QY 321 CAAAACCAACCAAAAAAACAACATA 352
DB 57650 CAAAACCAACCAAAAAAACAACATA 57619

RESULT 9
US-09-949-016-12683/C
; Sequence 12683, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; FILE REFERENCE: WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
; FILE REFERENCE: CLO01307
; CURRENT APPLICATION NUMBER: US/09/949,016
; PRIOR FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 12683
; LENGTH: 152524
; TYPE: DNA
; ORGANISM: Human
US-09-949-016-12683

Query Match 38.4%; Score 153.4; DB 3; Length 152524;
Best Local Similarity 69.8%; Pred. No. 7.9e-35;
Matches 224; Conservative 1; Mismatches 87; Indels 9; Gaps 1;

QY 36 CAAATATTAATTAAGCATTTGTCAGGCCAGGATGACATCGGCTGAATGCTGTATCCCA 95
DB 23848 CTAATTCCTCAAAATTAATTTCTTAATCTCTGCGGGGCGAGTGCCTCAAGCTGTATCCCA 23789
QY 96 GCACCTCGGAGGCCAAGTGGCGGATCACTGAGTCAAGAGATCGAGACCATCTCTGG 155
DB 23788 GCACCTTGGAGGCTGAGTGGGTGATCACTGAGTCAAGAGATCGAGACCATCTCTGG 23729
QY 156 CCAACATGATGAAACCCCGTCTTTTCTTAAATTAATCAAAAAATAGTGGCGATGTTGGCAC 215
DB 23728 CCAATATGATGAAACCCAGTCTCTAATAAATAAATAAATAAATAGCCGGCGTGGTGGCAT 23669
QY 216 ACACCTGATGCCAGCTACTCAGGAGCGGAGATTGAGTGAAGTGAATCGCAGAGTG 275
DB 23668 ATGCTGTAGTCCAGCTACTCAGGAGCGTGAAGGCTGAAGGCTGAAGGCTGAAGGCTGAAGG 23609
QY 276 AGCCGAATATCAAGATCAAGAGTGAAG-----CAGAGTGAAGACCCGCTTCAAAAA 326
DB 23608 CGAAAGTTGAGTGAAGCGGAGGTGTGCGCACTGCGACAGAGAGAGACTGTCTCAAAAA 23549

QY 327 CAACACCAAAAAACAACAAAA 347
DB 23548 CAACACCAAAAAACAACAAAA 23528

RESULT 10
US-09-949-016-13194/C
; Sequence 13194, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; FILE REFERENCE: WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
; FILE REFERENCE: CLO01307
; CURRENT APPLICATION NUMBER: US/09/949,016
; PRIOR FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 13194
; LENGTH: 152524
; TYPE: DNA
; ORGANISM: Human
US-09-949-016-13194

Query Match 38.4%; Score 153.4; DB 3; Length 152524;
Best Local Similarity 69.8%; Pred. No. 7.9e-35;
Matches 224; Conservative 1; Mismatches 87; Indels 9; Gaps 1;

QY 36 CAAATATTAATTAAGCATTTGTCAGGCCAGGATGACATCGGCTGAATGCTGTATCCCA 95
DB 23848 CTAATTCCTCAAAATTAATTTCTTAATCTCTGCGGGGCGAGTGCCTCAAGCTGTATCCCA 23789
QY 96 GCACCTCGGAGGCCAAGTGGCGGATCACTGAGTCAAGAGATCGAGACCATCTCTGG 155
DB 23788 GCACCTTGGAGGCTGAGTGGGTGATCACTGAGTCAAGAGATCGAGACCATCTCTGG 23729
QY 156 CCAACATGATGAAACCCCGTCTTTTCTTAAATTAATCAAAAAATAGTGGCGATGTTGGCAC 215
DB 23728 CCAATATGATGAAACCCAGTCTCTAATAAATAAATAAATAAATAGCCGGCGTGGTGGCAT 23669
QY 216 ACACCTGATGCCAGCTACTCAGGAGCGGAGATTGAGTGAAGTGAATCGCAGAGTG 275
DB 23668 ATGCTGTAGTCCAGCTACTCAGGAGCGTGAAGGCTGAAGGCTGAAGGCTGAAGGCTGAAGG 23609
QY 276 AGCCGAATATCAAGATCAAGAGTGAAG-----CAGAGTGAAGACCCGCTTCAAAAA 326
DB 23548 CAACACCAAAAAACAACAAAA 23528

RESULT 11
US-09-949-016-16011/C
; Sequence 16011, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; FILE REFERENCE: WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
; FILE REFERENCE: CLO01307
; CURRENT APPLICATION NUMBER: US/09/949,016
; PRIOR FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20

PRIOR APPLICATION NUMBER: 60/237,768
PRIOR FILING DATE: 2000-10-03
PRIOR APPLICATION NUMBER: 60/231,498
PRIOR FILING DATE: 2000-09-08
NUMBER OF SEQ ID NOS: 207012
SOFTWARE: FASTSEQ for Windows Version 4.0
SEQ ID NO 16011
LENGTH: 40091
TYPE: DNA
ORGANISM: Human
US-09-949-016-16011

Query Match 38.2%; Score 152.8; DB 3; Length 40091;
Best Local Similarity 75.0%; Pred. No. 6.9e-35;
Matches 219; Conservative 1; Mismatches 63; Indels 9; Gaps 2;

QY 84 CCTGTAATCCGAGCACTTCGGAGGCGCAAGGTGGCGGATCAGCTGAGTCAAGATCG 143
DB 30035 CCTAATATCCGAGCACTTTGGAGGCGGAGGCAAGTATCATTGAGGCGAGAGTCCG 29976
QY 144 AGACCATCTGGCCCAACATGTTAAACCCCGTCTTTACTTAAATAACAAAAATAGCTGG 203
DB 29975 AGACCATCTGGCCCAACATGTTAAATCCATCTCTTAAATAACAAAAATTAAGCTGG 29916
QY 204 GCATGTGGGACACACCTGTATGCCAGCTACTCAGAGGCGGAGATTGAGTGAAGCTGA 263
DB 29915 GCATGTGGGTCACAGCCTGTATATCCAGCTACTTGGAGGCTGAGATTGAGTGAAGCTGA 29856
QY 264 GATCGAGAGTGAAGCCGAATACAGATCAGAGTGAAGAGTGAAGACGCGTCTCA 323
DB 29855 GATCG-----TGCCACTGCACTGAGAGCTG--GGCAACAGATGAAGATCCATCTCA 29805
QY 324 AAACCAACACAAAAAACAACCAATTAAGATTTGCTGCGGTTC 375
DB 29804 AAACCAATTAACAAAAACAGCAACAAACAACTTAAGTTTCAAGGTTC 29753

RESULT 12

US-09-949-016-69587/C
Sequence 69587, Application US/09949016
Patent No. 6812339
GENERAL INFORMATION:
APPLICANT: VENTER, J. Craig et al.
TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
TITLE OF INVENTION: WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
FILE REFERENCE: C1.001307
CURRENT APPLICATION NUMBER: US/09/949,016
CURRENT FILING DATE: 2000-04-14
PRIOR APPLICATION NUMBER: 60/241,755
PRIOR FILING DATE: 2000-10-20
PRIOR APPLICATION NUMBER: 60/237,768
PRIOR FILING DATE: 2000-10-03
PRIOR APPLICATION NUMBER: 60/231,498
PRIOR FILING DATE: 2000-09-08
NUMBER OF SEQ ID NOS: 207012
SOFTWARE: FASTSEQ for Windows Version 4.0
SEQ ID NO 69587
LENGTH: 601
TYPE: DNA
ORGANISM: Human
US-09-949-016-69587

Query Match 38.0%; Score 152; DB 3; Length 601;
Best Local Similarity 74.0%; Pred. No. 2.2e-35;
Matches 225; Conservative 2; Mismatches 57; Indels 20; Gaps 2;

QY 81 ATGCTGTATATCCGAGCACTTCGGAGGCGCAAGGTGGCGGATCAGCTGAGTCAAGAGA 140
DB 487 AGCGCTGTATATCCGAGCACTTTGGAGGCGTAAAGTGGCGGATCAGCTGAGTCAAGAGT 428
QY 141 TCGAGACCATCTGGCCCAACATGTTAAACCCCGTCTTTACTTAAATAACAAAAATAGC 200
DB 427 TCGAGACCATCTGGCCCAACATGTTAAACCCCGTCTTTACTTAAATAACAAAAATAGC 368

QY 201 TGGGCAATGTGGGACACACCTGTATGTCAGACTACTGAGAGCC-----GGAGATTGAGT 256
DB 367 CGGACATGTGTGGGACGCACTGTATATCCAGCTACTTGGAGGCTTGGAGGAGATCACTT 308
QY 257 GAGCTGATGTGGGAGTGGAGCCGAATTCACAGATCA-----CAGAGTG 300
DB 307 GAACCCGAGGAGGAGAGTGTTCAGTGAAGCCAGATCAAGCACTGACCTGAGCTGGGT 248
QY 301 AGCAGGTGAGACGCGCTCTCAAAAACAACAACAAAAAACAACAAAAAATTAAGCATTTG 360
DB 247 GACAGAGTGAAGACTCGCTCTCAAAAACAACAAAAAACAACAAAAAACAACAAAAA 188
QY 361 TCCA 364
DB 187 ACCA 184

RESULT 13

US-09-949-016-12784/C
Sequence 12784, Application US/09949016
Patent No. 6812339
GENERAL INFORMATION:
APPLICANT: VENTER, J. Craig et al.
TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
TITLE OF INVENTION: WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
FILE REFERENCE: C1.001307
CURRENT APPLICATION NUMBER: US/09/949,016
CURRENT FILING DATE: 2000-04-14
PRIOR APPLICATION NUMBER: 60/241,755
PRIOR FILING DATE: 2000-10-20
PRIOR APPLICATION NUMBER: 60/237,768
PRIOR FILING DATE: 2000-10-03
PRIOR APPLICATION NUMBER: 60/231,498
PRIOR FILING DATE: 2000-09-08
NUMBER OF SEQ ID NOS: 207012
SOFTWARE: FASTSEQ for Windows Version 4.0
SEQ ID NO 12784
LENGTH: 17050
TYPE: DNA
ORGANISM: Human
FEATURE:
NAME/KEY: misc feature
LOCATION: (1)---(17050)
OTHER INFORMATION: n = A,T,C or G
US-09-949-016-12784

Query Match 37.9%; Score 151.4; DB 3; Length 17050;
Best Local Similarity 76.4%; Pred. No. 1.3e-34;
Matches 201; Conservative 1; Mismatches 52; Indels 9; Gaps 1;

QY 81 ATGCTGTATATCCGAGCACTTCGGAGGCGCAAGGTGGCGGATCAGCTGAGTCAAGAGA 140
DB 11937 AAGCTGTATATCCGAGCACTTTGGAGGCGGAGGCGGATCACTTGAAGTCAAGAGT 11878
QY 141 TCGAGACCATCTGGCCCAACATGTTAAACCCCGTCTTTACTTAAATAACAAAAATAGC 200
DB 11877 TCGAGACCATCTGGCCCAACATGTTAAACCTCGTCTTACTTAAATAACAAAAATAGC 11818
QY 201 TGGGCAATGTGGGACACACCTGTATGTCAGACTACTCAGAGGCGGAGATTGAGTGAAG 260
DB 11817 CGGACATGTGTGGGCGCGGCTGTATATCCAGCTACTCGGAGGCGGAGTGGAGCGAGC 11758
QY 261 TGAATGTGAGAGTGAAGCGGAAATCAAGATCAAGATGAGCAGAGTGAAGACCGCTCT 320
DB 11757 AGAGATGTGCTGATTCAGCTCAGTCTGGGGA-----CAAGGAGAGCTTCGCTCT 11707
QY 321 CAACCAACCAACCAACCAACCAACCAACCAACCAACCAACCAACCAACCAACCAACCAAC 343
DB 11706 CAACCAACCAACCAACCAACCAACCAACCAACCAACCAACCAACCAACCAACCAACCAAC 11684

RESULT 14


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US-09-949-016-13680/c
; Sequence 13680, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
; FILE REFERENCE: CL001307
; CURRENT APPLICATION NUMBER: US/09/949,016
; CURRENT FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 13680
; LENGTH: 17050
; TYPE: DNA
; ORGANISM: Human
; FEATURE:
; NAME/KEY: misc.feature
; LOCATION: (1)..(17050)
; OTHER INFORMATION: n = A,T,C or G
US-09-949-016-13680

Query Match          37.9%; Score 151.4; DB 3; Length 17050;
Best Local Similarity 76.4%; Pred. No. 1.3e-34;
Matches 201; Conservative 1; Mismatches 52; Indels 9; Gaps 1;

QY      81 ATGCTGTAAATCCCAAGCACTTCGGAGAGCCCAAGTGGCGGATCACTGAGGTCAAGAGA 140
DB      11937 AAGCTGTAAATCCCAAGCACTTCGGAGAGCCCAAGTGGCGGATCACTGAGGTCAAGAGA 11878

QY      141 TCGAGACCATCTGGCGCAACATGATGAAACCCCGTCTTACTTAAATAACAAATAATAGC 200
DB      11877 TCGAGACCATCTGGCGCAACATGATGAAACCCCGTCTTACTTAAATAACAAATAATAGC 11818

QY      201 TGGGATGATGGGACACACACTGTAGTCCAGCTACTCAGAGCCGAGATTGCAAGTGAAC 260
DB      11817 CGCAGTGTGGCGCGCGCGCTGTATCCAGCTACTCAGAGCCGAGATTGCAAGTGAAC 11758

QY      261 TGAAGTCCAGAGTGAAGCCGAAATACAGATCAAGATGAGAGTGAAGAGCCGCTCT 320
DB      11757 AGAGATCGTGCATTCAGCTCCAGTCTGGGCGA-----CACAGGAGAGCTCCGCTCT 11707

QY      321 CAAAAGCAACAACAAAAACAA 343
DB      11706 CAAAAGCAACAACAAAAACAA 11684

RESULT 15
US-09-949-016-13784
; Sequence 13784, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
; FILE REFERENCE: CL001307
; CURRENT APPLICATION NUMBER: US/09/949,016
; CURRENT FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 13784

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; LENGTH: 20099
; TYPE: DNA
; ORGANISM: Human
; FEATURE:
; NAME/KEY: misc.feature
; LOCATION: (1)..(20099)
; OTHER INFORMATION: n = A,T,C or G
US-09-949-016-13784

Query Match          37.7%; Score 150.8; DB 3; Length 20099;
Best Local Similarity 74.0%; Pred. No. 2.1e-34;
Matches 225; Conservative 1; Mismatches 58; Indels 20; Gaps 2;

QY      81 ATGCTGTAAATCCCAAGCACTTCGGAGAGCCCAAGTGGCGGATCACTGAGGTCAAGAGA 140
DB      8799 AAGCTGTAAATCCCAAGCACTTCGGAGAGCCCAAGTGGCGGATCACTGAGGTCAAGAGA 8858

QY      141 TCGAGACCATCTGGCGCAACATGATGAAACCCCGTCTTACTTAAATAACAAATAATAGC 200
DB      8859 TCGAGACCATCTGGCGCAACATGATGAAACCCCGTCTTACTTAAATAACAAATAATAGC 8918

QY      201 TGGGATGATGGGACACACACTGTAGTCCAGCTACTCAGAGCC---GAGATTGCAAGT 256
DB      8919 CGGACATGATGGGACACACACTGTAGTCCAGCTACTTGGAGGCTGAGGAGATCACTT 8978

QY      257 GAGCTGAATCGCAGAGTGAAGCCGAAATCAAGATCA-----CAGAGTG 300
DB      8979 GAACCCGGAGGAGGAGTTCAGTGAAGCCCAAGATCAAGATCACTGACCTGAGT 9038

QY      301 AGCAGATGAGAGCCCGCTCAAAAACAAACAAACAAACAAACAAACAAACAAACAAACAAAC 360
DB      9039 GACAGATGAGAGCTCGTCTCAAAAACAAACAAACAAACAAACAAACAAACAAACAAAC 9098

QY      361 TCCA 364
DB      9099 ACCA 9102

RESULT 16
US-09-949-016-12096/c
; Sequence 12096, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
; FILE REFERENCE: CL001307
; CURRENT APPLICATION NUMBER: US/09/949,016
; CURRENT FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 12096
; LENGTH: 19566
; TYPE: DNA
; ORGANISM: Human
US-09-949-016-12096

Query Match          37.6%; Score 150.4; DB 3; Length 19566;
Best Local Similarity 72.1%; Pred. No. 2.7e-34;
Matches 196; Conservative 0; Mismatches 76; Indels 0; Gaps 0;

QY      81 ATGCTGTAAATCCCAAGCACTTCGGAGAGCCCAAGTGGCGGATCACTGAGGTCAAGAGA 140
DB      14763 ATGCTGTAAATCCCAAGCACTTCGGAGAGCCCAAGTGGCGGATCACTGAGGTCAAGAGA 14704

QY      141 TCGAGACCATCTGGCGCAACATGATGAAACCCCGTCTTACTTAAATAACAAATAATAGC 200

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Db 14703 TCAAGACCGAGCTGGCCCAACATGTTGTAACCCCGTCTCTACTATAAAATACAAAAATTATG 14644
Qy 201 TGGGCAATGTTGGACACACCTGTAGTCCAGCTACTACAGAGCCGGAGATTGCGTGAAC 260
Db 14643 TGGGCAATGTTGGACACCTGTAGTCCAGCTACTACAGAGCCGGAGATTGCGTGAAC 14584
Qy 261 TGAAGTCAGAGTGAAGCGGAATACAGATCACAGATGAGAGTGAAGACCKCGCT 320
Db 14583 CTGTAACCCAGAGTGAAGTGTGATGAGCCGAGATTGTGCACTTCATCTCA 14524
Qy 321 CAAAAACAACAACAAAAACCAATACATA 352
Db 14523 AAAAACTTCACTTCAAAAAAAGAGAAA 14492

RESULT 17
US-09-949-016-14114/c
; Sequence 14114, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; FILE REFERENCE: CL001307
; CURRENT FILING DATE: 2000-04-14
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 14114
; LENGTH: 19567
; TYPE: DNA
; ORGANISM: Human
US-09-949-016-14114

Query Match 37.6%; Score 150.4; DB 3; Length 19567;
Best Local Similarity 72.1%; Pred. No. 2.7e-34;
Matches 196; Conservative 0; Mismatches 76; Indels 0; Gaps 0;

Qy 81 ATGCTGTAAATCCAGCACTTGGGAGCCGCAAGTGGCGGATTCACCTGATCAAGCA 140
Db 14763 ATGCTGTAAATCCAGCACTTGGGAGCCGCAAGTGGCGGATTCACCTGATCAAGCA 14704
Qy 141 TCGAGACCATCTGGCCCAACATGTGAAACCCGCTCTTTACTTAAATAACAAAAATAGC 200
Db 14703 TCAAGACCGAGCTGGCCCAACATGTGAAACCCGCTCTTTACTTAAATAACAAAAATAGC 14644
Qy 201 TGGGCAATGTTGGACACACCTGTAGTCCAGCTACTACAGAGCCGGAGATTGCGTGAAC 260
Db 14643 TGGGCAATGTTGGACACACCTGTAGTCCAGCTACTACAGAGCCGGAGATTGCGTGAAC 14584
Qy 261 TGAAGTCAGAGTGAAGCGGAATACAGATCACAGATGAGAGTGAAGACCKCGCT 320
Db 14583 CTGTAACCCAGAGTGAAGTGTGATGAGCCGAGATTGTGCACTTCATCTCA 14524
Qy 321 CAAAAACAACAACAAAAACCAATACATA 352
Db 14523 AAAAACTTCACTTCAAAAAAAGAGAAA 14492

RESULT 18
US-09-949-016-14157
; Sequence 14157, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; FILE REFERENCE: CL001307
; CURRENT FILING DATE: 2000-04-14
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 14157
; LENGTH: 455726
; TYPE: DNA
; ORGANISM: Human
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: (1)..(455726)
; OTHER INFORMATION: n = A,T,C or G
US-09-949-016-14157

Query Match 37.4%; Score 149.6; DB 3; Length 455726;
Best Local Similarity 73.5%; Pred. No. 1.6e-33;
Matches 219; Conservative 1; Mismatches 69; Indels 9; Gaps 2;
```

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FILE REFERENCE: CL001307
; CURRENT APPLICATION NUMBER: US/09/949,016
; CURRENT FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 14157
; LENGTH: 455726
; TYPE: DNA
; ORGANISM: Human
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: (1)..(455726)
; OTHER INFORMATION: n = A,T,C or G
US-09-949-016-14157

Query Match 37.4%; Score 149.6; DB 3; Length 455726;
Best Local Similarity 73.5%; Pred. No. 1.6e-33;
Matches 219; Conservative 1; Mismatches 69; Indels 9; Gaps 2;

Qy 81 ATGCTGTAAATCCAGCACTTGGGAGCCGCAAGTGGCGGATTCACCTGATCAAGCA 140
Db 14383 ATGCTGTAAATCCAGCACTTGGGAGCCGCAAGTGGCGGATTCACCTGATCAAGCA 143892
Qy 141 TCGAGACCATCTGGCCCAACATGTGAAACCCGCTCTTTACTTAAATAACAAAAATAGC 200
Db 143893 TCAAGACCGAGTGGCCCAACATGTGAAACCCGCTCTTTACTTAAATAACAAAAATAGC 143952
Qy 201 TGGGCAATGTTGGACACACCTGTAGTCCAGCTACTACAGAGCCGGAGATTGCGTGAAC 260
Db 143953 TGGGCAATGTTGGACACACCTGTAGTCCAGCTACTACAGAGCCGGAGATTGCGTGAAC 144012
Qy 261 TGAAGTCAGAGTGAAGCGGAATACAGATCACAGATGAGAGTGAAGACCKCGCT 320
Db 144013 TGAAGTC-----TGCACCTGACATCCACCTCGGTGA-CAGAGTGAACATTTGCT 144063
Qy 321 CAAAAACAACAACAAAAACCAATACATA 378
Db 144064 TTAACAACAACAAAAAAGAGAAAATTCTACTTCTTGAATCCAGTGGCCCTG 144121

RESULT 19
US-09-949-016-11940
; Sequence 11940, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; FILE REFERENCE: CL001307
; CURRENT FILING DATE: 2000-04-14
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 11940
; LENGTH: 481115
; TYPE: DNA
; ORGANISM: Human
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: (1)..(481115)
; OTHER INFORMATION: n = A,T,C or G
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PRIOR APPLICATION NUMBER: 60/241,755
PRIOR FILING DATE: 2000-10-20
PRIOR APPLICATION NUMBER: 60/237,768
PRIOR FILING DATE: 2000-10-03
PRIOR APPLICATION NUMBER: 60/231,498
PRIOR FILING DATE: 2000-09-08
NUMBER OF SEQ ID NOS: 207012
SOFTWARE: FastSeq for Windows Version 4.0
SEQ ID NO 12517
LENGTH: 77994
TYPE: DNA
ORGANISM: Human
US-09-949-016-12517

Query Match 37.2%; Score 148.8; DB 3; Length 77994;
Best Local Similarity 72.7%; Pred. No. 1.4e-33;
Matches 192; Conservative 0; Mismatches 72; Indels 0; Gaps 0;

84 CCTGTAATCCAGCAGCTTGGGAGGCGCAAGGTGGCGGATGACCTGAGGTCAAGAGATCG 143
73058 CCTGTAATCCAGCAGCTTGGGAGGCGCAAGGTGGCGGATGACCTGAGGTCAAGAGATCG 72999

144 AGACCATCTGCGCCACATGTGTAAACCCGCTTTACTTAAATAACAAAAATAGCTGG 203
72998 AGACCATCTGCGCCACATGTGTAAACCCGCTTTACTTAAATAACAAAAATAGCTGG 72939

204 GCATGTGGCAGACACCTGTGTAGTCCCACTACTACAGAGCCGGAGATTGCAAGCTGA 263
72938 GCGGTGTGGCCACATCTGTATCCCACTACTTGGGAGGCTGAGGCAAGAAATTGCTAT 72879

264 GATGCGAGAGTGAGCGCAATATCACAGATCACAGATGAGAGAGAGACACCGCTCTCAA 323
72878 GAACCCAGAGGCGGAGGTTGCACTGCACTTACTAGCTTACAGCAAGAGCAACTGTCTC 72819

324 AAACACACACAAAAAACAACAAAAA 347
72818 AAACACACACAAAAAACAACAAAAA 72795

RESULT 23
US-09-949-016-16021/c
Sequence 16021, Application US/09949016
Patent No. 6812339
GENERAL INFORMATION:
APPLICANT: VENTER, J. Craig et al.
TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
FILE REFERENCE: CL001307
CURRENT APPLICATION NUMBER: US/09/949,016
PRIOR FILING DATE: 2000-04-14
PRIOR APPLICATION NUMBER: 60/241,755
PRIOR FILING DATE: 2000-10-20
PRIOR APPLICATION NUMBER: 60/237,768
PRIOR FILING DATE: 2000-10-03
PRIOR APPLICATION NUMBER: 60/231,498
PRIOR FILING DATE: 2000-09-08
NUMBER OF SEQ ID NOS: 207012
SOFTWARE: FastSeq for Windows Version 4.0
SEQ ID NO 16021
LENGTH: 77994
TYPE: DNA
ORGANISM: Human
US-09-949-016-16021

Query Match 37.2%; Score 148.8; DB 3; Length 77994;
Best Local Similarity 72.7%; Pred. No. 1.4e-33;
Matches 192; Conservative 0; Mismatches 72; Indels 0; Gaps 0;

84 CCTGTAATCCAGCAGCTTGGGAGGCGCAAGGTGGCGGATGACCTGAGGTCAAGAGATCG 143
73058 CCTGTAATCCAGCAGCTTGGGAGGCGCAAGGTGGCGGATGACCTGAGGTCAAGAGATCG 72999

144 AGACCATCTGCGCCACATGTGTAAACCCGCTTTACTTAAATAACAAAAATAGCTGG 203

72998 AGACCATCTGAGCTTAACATGATGAAACCCGCTTTACTTAAATAACAAAAATAGCTGG 72939

204 GCATGTGGCAGACACCTGTGTAGTCCCACTACTACAGAGCCGGAGATTGCAAGCTGA 263
72938 GCGGTGTGGCCACATCTGTATCCCACTACTTGGGAGGCTGAGGCAAGAAATTGCTAT 72879

264 GATGCGAGAGTGAGCGCAATATCACAGATCACAGATGAGAGAGAGACACCGCTCTCAA 323
72878 GAACCCAGAGGCGGAGGTTGCACTGCACTTACTAGCTTACAGCAAGAGCAACTGTCTC 72819

324 AAACACACACAAAAAACAACAAAAA 347
72818 AAACACACACAAAAAACAACAAAAA 72795

RESULT 24
US-09-949-016-13460/c
Sequence 13460, Application US/09949016
Patent No. 6812339
GENERAL INFORMATION:
APPLICANT: VENTER, J. Craig et al.
TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
FILE REFERENCE: CL001307
CURRENT APPLICATION NUMBER: US/09/949,016
PRIOR FILING DATE: 2000-04-14
PRIOR APPLICATION NUMBER: 60/241,755
PRIOR FILING DATE: 2000-10-20
PRIOR APPLICATION NUMBER: 60/237,768
PRIOR FILING DATE: 2000-10-03
PRIOR APPLICATION NUMBER: 60/231,498
PRIOR FILING DATE: 2000-09-08
NUMBER OF SEQ ID NOS: 207012
SOFTWARE: FastSeq for Windows Version 4.0
SEQ ID NO 13460
LENGTH: 8100
TYPE: DNA
ORGANISM: Human
US-09-949-016-13460

Query Match 37.2%; Score 148.6; DB 3; Length 8100;
Best Local Similarity 74.9%; Pred. No. 6.4e-34;
Matches 200; Conservative 1; Mismatches 60; Indels 6; Gaps 1;

83 GCTGTATATCCAGCAGCTTGGGAGGCGCAAGGTGGCGGATGACCTGAGGTCAAGATC 142
6648 GCTGTATATCCAGCAGCTTGGGAGGCGCAAGGTGGGATGATCAGGTGAGGTTC 6589

143 GAGACCATCTGCGCCACATGTGTAAACCCGCTTTACTTAAATAACAAAAATAGCTGG 202
6588 AAGACCATCTGCGCCACATGTGTAAACCCGCTTTACTTAAATAACAAAAATAGCTGG 6529

203 GCATGTGGCAGACACCTGTGTAGTCCCACTACTACAGAGCCGGAGATTGCAAGCTG 262
6528 GGCATGTGGGAGGCGCTGTATCCCACTACTTGGGAGGCTGAGGCAAGAAATCGCT 6469

263 AGATGCGAGATGAGCGCAATATCACAGATC-----ACAGATGAGCAGATGAGACGCC 316
6468 TGAACCCAGAGGAGAGAGGTTGCACTGAGCACTCAACTGGGCAACAGAGTGAAGCTCT 6409

317 GTCTCAAAAAACAACAAAAAACA 343
6408 GTCTCAAAAAAACAACAAAAAACA 6382

RESULT 25
US-09-949-016-13461/c
Sequence 13461, Application US/09949016
Patent No. 6812339
GENERAL INFORMATION:
APPLICANT: VENTER, J. Craig et al.
TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED

;; TITLE OF INVENTION: WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
;; FILE REFERENCE: CL001307
;; CURRENT APPLICATION NUMBER: US/09/949,016
;; CURRENT FILING DATE: 2000-04-14
;; PRIOR APPLICATION NUMBER: 60/241,755
;; PRIOR FILING DATE: 2000-10-20
;; PRIOR APPLICATION NUMBER: 60/237,768
;; PRIOR FILING DATE: 2000-10-03
;; PRIOR APPLICATION NUMBER: 60/231,498
;; PRIOR FILING DATE: 2000-09-08
;; NUMBER OF SEQ ID NOS: 207012
;; SOFTWARE: FastSeq for Windows Version 4.0
;; SEQ ID NO 13461
;; LENGTH: 8100
;; TYPE: DNA
;; ORGANISM: Human
US-09-949-016-13461

Query Match 37.2%; Score 148.6; DB 3; Length 8100;
Best Local Similarity 74.9%; Pred. No. 6.4e-34;
Matches 200; Conservative 1; Mismatches 60; Indels 6; Gaps 1;

QY 83 GCCTTAATCCGACACTTCCGAGAGCCCAAGGTGGCGGATCACTGAGTCAAGATC 142
DB 6648 GCCTTAATCCGACACTTGGGAGGCGCAAGGTAGTGATCACTGAGTCAAGGTTCC 6589

QY 143 GAGACCATCTGGCCCAACATGGTGAACCCGCTTTACTAAATATCAAAATATGCTG 202
DB 6588 AAGACCACTTGGCCCAACATGGTGAACCCGCTTTACTAAATATCAAAATATGCTG 6529

QY 203 GGCATGTTGGGACACACCTGTAGTCCAGCTACTCAGAGCCGAGATTGACAGCTG 262
DB 6528 GGCATGTTGGGACGCGGCGCTGTATCCAGCTACTTGGAGGCTGAGGAGGATTCCT 6469

QY 263 AGATGCGAGTGCAGCCGAAATCAGATC-----ACAGAGTGCAGAGTGCAGACGCC 316
DB 6468 TGAACCCGAGGAGGAGAGTTCAGTGCAGCTCCAACTGGGCAACAGAGTGCAGCTCT 6409

QY 317 GTCTCAAAAACACAAACAAAACAA 343
DB 6408 GTCTCAAAAACAAAACAAAACAA 6382

RESULT 26
US-09-949-016-152016
; Sequence 152016, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
; FILE REFERENCE: CL001307
; CURRENT APPLICATION NUMBER: US/09/949,016
; CURRENT FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 152016
; LENGTH: 601
; TYPE: DNA
; ORGANISM: Human
US-09-949-016-152016

Query Match 37.0%; Score 148; DB 3; Length 601;
Best Local Similarity 75.0%; Pred. No. 3.4e-34;
Matches 213; Conservative 1; Mismatches 61; Indels 9; Gaps 2;

QY 84 CCTGTAAATCCGACACTTCCGAGAGCCCAAGGTGGCGGATCACTGAGTCAAGATCG 143

DB 326 CCTATATCCGACGACTTGGGAGGCCGAGGAGGTAGATCACTGAGCCGAGAGTCCG 385
QY 144 AGACCATCTGGCCCAACATGTGTAAACCCGCTTTACTAAATATCAAAATATGCTG 203
DB 386 AGACCACTGCGCCCAACATGTGTAAATCCATCTTACTAAACAAATATGCTG 445

QY 204 GCATGTTGGGACACACCTGTAGTCCAGCTACTCAGAGCCGAGATTGCAGTGAGCTGA 263
DB 446 GCATGTTGGGACGCGCTGTATCCAGCTACTTGGGAGGCTGAGATTGCAGTGAGCTGA 505

QY 264 GATGCGAGAGTGCAGCCGAAATCAGATCAGAGAGTGCAGAGTGCAGACGCCGCTCA 323
DB 506 GATG-----TGCACTGCATGCGAGACTG- GCGAAGAGATGAATCTCACTCA 556

QY 324 AAACACACACAAACAAACAAACAAACAAACAAACAACTTGTCT 367
DB 557 AAAAAATATAAAAAACAAACAAACAAACAAACAACTTGTCT 600

RESULT 27
US-09-949-002-2953/c
; Sequence 2953, Application US/09949002
; Patent No. 6900016
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; WITH INFLAMMATORY AUTOIMMUNE DISEASE, METHODS OF DETECTION
; FILE REFERENCE: CL000790
; CURRENT APPLICATION NUMBER: US/09/949,002
; CURRENT FILING DATE: 2000-01-28
; PRIOR APPLICATION NUMBER: 60/231,401
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 10823
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 2953
; LENGTH: 601
; TYPE: DNA
; ORGANISM: Human
US-09-949-002-2953

Query Match 37.0%; Score 147.8; DB 3; Length 601;
Best Local Similarity 76.9%; Pred. No. 3.9e-34;
Matches 176; Conservative 0; Mismatches 47; Indels 0; Gaps 0;

QY 81 ATGCTGTAAATCCGACGACTTCCGAGAGCCCAAGGTGGCGGATCACTGAGTCAAGAGA 140
DB 232 ATGCTGTAAATCCGACGACTTGGGAGGCCGAGGTTGGGTGATGCCCTGAGTCAAGAGT 173

QY 141 TCGAGACCATCTGGCCCAACATGTGTAAACCCGCTTTACTAAATATCAAAATATGAC 200
DB 172 TTGAGACCCGCGCTGGCCCAACATGTGTAAACCCGCTACTACTAAATATCAAAATATGAC 113

QY 201 TGGGATGTTGGGACACACCTGTAGTCCAGCTACTCAGAGCCGAGATTGCAGTGAGC 260
DB 112 TGGGATGTTGGGACACACCTGTAGTCCAGCTACTCAGAGGCTGAGGAGATGGAATTG 53

QY 261 TGAATGCGAGAGTGCAGCCGAAATCAGATCAGAGTGCAGAGTGCAGC 303
DB 52 CTGTAACCCAGAGGTGAGAGTTGCAATGAGCCCAAGATGATGTC 10

RESULT 28
US-09-949-002-2954/c
; Sequence 2954, Application US/09949002
; Patent No. 6900016
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; WITH INFLAMMATORY AUTOIMMUNE DISEASE, METHODS OF DETECTION
; FILE REFERENCE: CL000790

```

: CURRENT APPLICATION NUMBER: US/09/949,002
: CURRENT FILING DATE: 2000-01-28
: PRIOR APPLICATION NUMBER: 60/231,401
: PRIOR FILING DATE: 2000-09-08
: NUMBER OF SEQ ID NOS: 10823
: SOFTWARE: FASTSEQ for Windows Version 4.0
: SEQ ID NO 2954
:
: LENGTH: 601
:
: TYPE: DNA
:
: ORGANISM: Human
:
: US-09-949-002-2954

```

Query Match	37.0%	Score 147.8	DB 3	Length 601
Best Local Similarity	76.9%	Pred.No. 3.9e-34		
Matches 176	Conservative	0	Mismatches 47	Indels 0
				Gaps 0

QY	81	TTGCGCTGTAATCCGAGCACTTGGGAGGGCGAAGTGGGGGAACTACCTAGAGGCAAGAGA	140
Db	253	ATGCTCTGTAATCCAGACACTTTGGGAGGGCGAAGTGGGATGCGCTGAGGTCAAGAGT	194
QY	141	TCGAGACCATCCTGGCGCAACATGATGTAACCCCGCTTTACTTAAAAATCAAAAAATAGC	200
Db	193	TTGATACCGGCTTGGCGAACATGATGTAACCCCGTCACTACTTAAAAATCAAAAAATTAGC	134
QY	201	TGGGCATGCTGGCACACACTGTAGTCCAGCTTCAAGAGCGCGAGATTGCAAGTGAGC	260
Db	133	TGGGCATGCTGGCACACACTGTAGTCCAGCTTCAAGAGCGCTGAGGCATGAGATTG	74
QY	261	TGAGATCGCAGAGTGAGCCGAATCAACAATCAACAAGTGAGC	303
Db	73	CTTGAACCCAGAGGTGAGGTTGACGTAGAGCCAAATGCTGC	31

```

RESULT 29
US-09-949-002-4465/C
; Sequence 4465, Application US/09949002
; Patent No. 6500016
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; TITLE OF INVENTION: WITH INFLAMMATORY AUTOIMMUNE DISEASE, METHODS OF DETECTION
; TITLE OF INVENTION: AND USES THEREOF
; FILE REFERENCE: CLO00790
; CURRENT APPLICATION NUMBER: US/09/949,002
; CURRENT FILING DATE: 2000-01-28
; PRIOR APPLICATION NUMBER: 60/231,401
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 10823
; SOFTWARE: FASTSEQ for Windows Version 4.0
; SEQ ID NO 4465
; LENGTH: 601
; TYPE: DNA
; ORGANISM: Human
; US-09-949-002-4465

```

Query Match	37.0%	Score 147.8	DB 3	Length 601
Best Local Similarity	78.9%	Pred. No. 3.9e-34		
Matches 176, Conservative	0	Mismatches 47	Indels 0	Gaps 0

QY 81 AAGCCTGTAATCCAGACACTTCGAGAGGCCAAGTGGCGGATCACCCTGAGTCAAGAGA 140
DB 232 ATGCCCTGTAATCCAGACACTTTGGGAGGCGGAAGTGGTGGATTCGCTTGAGTCAAGAGT 173
QY 141 TCGAGACCAATCCTGGGCCAACATGTGTAAACCCCGTCTTTACTTAAAAATACAAAAAATAGC 200
DB 172 TTGAGACCCGCGCTGGCCAAATGTGTAAACCCCGTCACTTAAAAATACAAAAAATTAGC 113
QY 201 TGGGCATGTGTGCACACACTGTAGTCCACGACTACTCAGAGCCGGGAGATTGCAGTAGC 26
DB 112 TGGGCATGTGTGCACACACTTGTAGTCCACGACTACTCAGAGGCTGGGAGATGCAGATTG 53
QY 261 TGGAGTGCAGAGTGAAGCCGAAATCAACAGTCAACAGTGAAGC 303

Db 52 CTGAACCCAGAAGGTGAGGTTGCAGTGAGCCAGAATGTTGC 10

```

RESULT 30
US-09-949-002-4466/C
; Sequence 4466, Application US/09949002
; Patent No. 690016
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; WITH INFLAMMATORY AUTOIMMUNE DISEASE, METHODS OF DETECTION
; TITLE OF INVENTION: AND USES THEREOF
; FILE REFERENCE: CL000790
; CURRENT APPLICATION NUMBER: US/09/949,002
; CURRENT FILING DATE: 2000-01-28
; PRIOR APPLICATION NUMBER: 60/231,401
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 10823
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 4466
; LENGTH: 601
; TYPE: DNA
; ORGANISM: Human
US-09-949-002-4466

```

Query Match	37.0%;	Score 147.8;	DB 3;	Length 601;
Best Local Similarity	78.9%;	Pred. No. 3.9e-34;		
Matches 176; Conservative	0;	Mismatches 47;	Indels 0;	Gaps 0;

QY 8 ATCCCTGTATCCCAAGCATTTGGGAGGCGCAAGGGGCGGATCACTGAGGTCAAGAGA 140
Db 253 ATGCTCTGTATCCCAAGCATTTGGGAGGCGCAAGGGGCGGATCACTGAGGTCAAGAGT 194
QY 141 TCGAGACCATCTCTGGCAACATGTGTAAACCCCGCTTTACTTAAATAATCAAAAAATAGC 200
Db 193 TTGAGACCGGCGCTGGCCAACTGTGTAAACCCCGCTCACTAATAAATAATTAATAGC 134
QY 201 TGGGCATGTGTGGACACACACTGTAGTCCCACTCAAGATCTCAGAGACCGGAGATTGCAGTGAGC 266
Db 133 TGGGCATGTGTGGACACACACTGTAGTCCCACTCAAGATCTCAGAGACCGGAGATTGCAAGATTG 74
QY 261 TGAGATCGCAGAGTGAAGCGGAAATACAGATCAGCAAGTAGAGC 303
Db 73 CTTGAACCCCAAGGTGTGAGTTTGCACTGTAGCCAAAGATGTGTC 31

```

1      RESULT 31
2      US-09-949-002-650
3      ; Sequence 650, Application US/09949002
4      ; Patent No. 6500016
5      ; GENERAL INFORMATION:
6      ; APPLICANT: VENTER, J. Craig et al.
7      ; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
8      ; TITLE OF INVENTION: WITH INFLAMMATORY AUTOIMMUNE DISEASE, METHODS OF DETECTION
9      ; TITLE OF INVENTION: AND USES THEREOF
10     ; FILE REFERENCE: CLO000790
11     ; CURRENT APPLICATION NUMBER: US/09/949,002
12     ; CURRENT FILING DATE: 2000-01-28
13     ; PRIOR APPLICATION NUMBER: 60/231,401
14     ; PRIOR FILING DATE: 2000-09-08
15     ; NUMBER OF SEQ ID NOS: 10823
16     ; SOFTWARE: FASTSEQ for Windows Version 4.0
17     ; SEQ ID NO 650
18     ; LENGTH: 60595
19     ; TYPE: DNA
20     ; ORGANISM: Human
21     ; US-09-949-002-650

```

Query Match	37.0%;	Score 147.8;	DB 3;	Length 60595;
Best Local Similarity	78.9%;	Pred. No. 2.5e-33;		
Matches 176; Conservative	0;	Mismatches 47;	Indels 0;	Gaps 0
QY	81	ATGCCTGTAATCCCAAGCACTTCGGGAGGCCAAAGTGCGCGGATCACTTGAGGTCAAGAGA	140	

81 ATGCCTGTAATCCAGCACTTCGGAGGCCAAGTGGCGGATCACCTGAGGTCAGAGA 140

DB 55522 ATGCGTGAATCCAGCACTTTGGAGGCGGAGGTGGGTGATCGCTGAGGTCAAGAGT 55581
QY 141 TGAAGACCATCTCTGGCCAAATGATGTAACCCCGTCTTACTTAAATAATCAAAAAATAGC 200
DB 55582 TTGAAGACCGGCTGGCCAAATGATGTAACCCCGTCTTACTTAAATAATCAAAAAATAGC 55641
QY 201 TGGGATGATGTCACACACTTGTAGTCCAGTACTCAGAGCCGGAATTCAGATGAGC 260
DB 55642 TGGGATGATGTCACACACTTGTAGTCCAGTACTCAGAGCCGGAATTCAGATGAGC 55701
QY 261 TGAGATCCGACAGTGGCCGAATATCAAGATCAAGATGAGC 303
DB 55702 CTTGAACCCAGAGGTGAGGTTGCAGTAGCCAAAGATGTGC 55744

RESULT 32
US-09-949-002-706
; Sequence 706, Application US/09949002
; Patent No. 6900016
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; TITLE OF INVENTION: WITH INFLAMMATORY AUTOIMMUNE DISEASE, METHODS OF DETECTION
; FILE REFERENCE: C1000790
; CURRENT APPLICATION NUMBER: US/09/949,002
; CURRENT FILING DATE: 2000-01-28
; PRIOR APPLICATION NUMBER: 60/231,401
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 10823
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 706
; LENGTH: 60595
; TYPE: DNA
; ORGANISM: Human
US-09-949-002-706

Query Match 37.0%; Score 147.8; DB 3; Length 60595;
Best Local Similarity 78.9%; Pred. No. 2.5e-33;
Matches 176; Conservative 0; Mismatches 47; Indels 0; Gaps 0;

QY 81 ATGCTGTAAATCCAGCACTTGGAGGCGGAGGTGGGTGATCGCTGAGGTCAAGAGA 140
DB 55522 ATGCTGTAAATCCAGCACTTGGAGGCGGAGGTGGGTGATCGCTGAGGTCAAGAGT 55581
QY 141 TGAAGACCATCTCTGGCCAAATGATGTAACCCCGTCTTACTTAAATAATCAAAAAATAGC 200
DB 55582 TTGAAGACCGGCTGGCCAAATGATGTAACCCCGTCTTACTTAAATAATCAAAAAATAGC 55641
QY 201 TGGGATGATGTCACACACTTGTAGTCCAGTACTCAGAGCCGGAATTCAGATGAGC 260
DB 55642 TGGGATGATGTCACACACTTGTAGTCCAGTACTCAGAGCCGGAATTCAGATGAGC 55701
QY 261 TGAGATCCGACAGTGGCCGAATATCAAGATCAAGATGAGC 303
DB 55702 CTTGAACCCAGAGGTGAGGTTGCAGTAGCCAAAGATGTGC 55744

RESULT 33
US-09-949-016-13849
; Sequence 13849, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; TITLE OF INVENTION: WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
; FILE REFERENCE: C1001307
; CURRENT APPLICATION NUMBER: US/09/949,016
; CURRENT FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768

; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 13849
; LENGTH: 86945
; TYPE: DNA
; ORGANISM: Human
; FEATURE:
; NAME/KEY: misc.feature
; LOCATION: (1) ... (86945)
; OTHER INFORMATION: n = A,T,C or G
US-09-949-016-13849

Query Match 37.0%; Score 147.8; DB 3; Length 86945;
Best Local Similarity 73.5%; Pred. No. 2.9e-33;
Matches 208; Conservative 1; Mismatches 58; Indels 16; Gaps 1;

QY 81 ATGCTGTAAATCCAGCACTTGGAGGCGGAGGTGGGTGATCGCTGAGGTCAAGAGA 140
DB 25003 ATGCTGTAAATCCAGCACTTGGAGGCGGAGGTGGGTGATCGCTGAGGTCAAGAGT 25062
QY 141 TGAAGACCATCTCTGGCCAAATGATGTAACCCCGTCTTACTTAAATAATCAAAAAATAGC 200
DB 25063 TTGAAGACCGGCTGGCCAAATGATGTAACCCCGTCTTACTTAAATAATCAAAAAATAGC 25122
QY 201 TGGGATGATGTCACACACTTGTAGTCCAGTACTCAGAGCCGGAATTCAGATGAGC 260
DB 25123 TGGGATGATGTCACACACTTGTAGTCCAGTACTCAGAGCCGGAATTCAGATGAGC 25182
QY 261 TGAGATCCGACAGTGGCCGAATATCAAGATCAAGATGAGC 304
DB 25183 CTGGAGGCGGAGGTGAGGTTGCAGTAGCCAAAGATGTGCATTCAGCTGGGCAACA 25242
QY 305 GAGTGAGACGCGCTCTCAAAAACAACAACAACAACAACAACAACAACAACAACAACA 347
DB 25243 GTGTGAGCTCCTCAATCTCAAAAACAACAACAACAACAACAACAACAACAACAACA 25285

RESULT 34
US-09-949-016-13850
; Sequence 13850, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; TITLE OF INVENTION: WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
; FILE REFERENCE: C1001307
; CURRENT APPLICATION NUMBER: US/09/949,016
; CURRENT FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03/231,498
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 13850
; LENGTH: 86945
; TYPE: DNA
; ORGANISM: Human
; FEATURE:
; NAME/KEY: misc.feature
; LOCATION: (1) ... (86945)
; OTHER INFORMATION: n = A,T,C or G
US-09-949-016-13850

Query Match 37.0%; Score 147.8; DB 3; Length 86945;
Best Local Similarity 73.5%; Pred. No. 2.9e-33;
Matches 208; Conservative 1; Mismatches 58; Indels 16; Gaps 1;

```
QY 81 ATGCTGTAATCCAGCACTTCGGAGGCCAAGTGGCGGATGACCTGAGTCAAGA 140
DB 25003 ATGCTGTAATCCAGCACTTCGGAGGCCAAGTGGCGGATGACCTGAGTCAAGA 25062
QY 141 TCGAGACCATCTGCGCCAACTGTGAACCCCGTCTTACTTAAATAACAAAAATAGC 200
DB 25063 TCGAGACCATCTGCGCCAACTGTGAACCCCGTCTTACTTAAATAACAAAAATAGC 25122
QY 201 TGGGATGTTGGCAACACCTGTAGTCCAGCTACTCAGAGCCGAGATTGCAGTAGC 260
DB 25123 TGGGATGTTGGCAACACCTGTAGTCCAGCTACTCAGAGCCGAGATTGCAGTAGC 25182
QY 261 TGAGATCCAGAGTGAAGCCGAATACAGAT-----CACAGAGTGAGCA 304
DB 25183 CTGGAGGCAAGAGTTGCAAGTGAAGTGAATGTCCTGACTCCAGCCTGGGCAACA 25242
QY 305 GAGTGAGACCKCGTCTCAAAAACAACAACAAAAA 347
DB 25243 GTGTGAGACTCTCATCTCAAAAAGAAATAAATAAATAAATAA 25285

RESULT 35
US-09-949-016-13851
; Sequence 13851, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; FILE REFERENCE: CL001307
; CURRENT APPLICATION NUMBER: US/09/949,016
; PRIOR FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 13851
; LENGTH: 86945
; TYPE: DNA
; ORGANISM: Human
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: (1)...(86945)
; OTHER INFORMATION: n = A,T,C or G
US-09-949-016-13851

Query Match 37.0%; Score 147.8; DB 3; Length 86945;
Best Local Similarity 73.5%; Pred. No. 2.9e-33;
Matches 208; Conservative 1; Mismatches 58; Indels 16; Gaps 1;

QY 81 ATGCTGTAATCCAGCACTTCGGAGGCCAAGTGGCGGATGACCTGAGTCAAGA 140
DB 25003 ATGCTGTAATCCAGCACTTCGGAGGCCAAGTGGCGGATGACCTGAGTCAAGA 25062
QY 141 TCGAGACCATCTGCGCCAACTGTGAACCCCGTCTTACTTAAATAACAAAAATAGC 200
DB 25063 TCGAGACCATCTGCGCCAACTGTGAACCCCGTCTTACTTAAATAACAAAAATAGC 25122
QY 201 TGGGATGTTGGCAACACCTGTAGTCCAGCTACTCAGAGCCGAGATTGCAGTAGC 260
DB 25123 TGGGATGTTGGCAACACCTGTAGTCCAGCTACTCAGAGCCGAGATTGCAGTAGC 25182
QY 261 TGAGATCCAGAGTGAAGCCGAATACAGAT-----CACAGAGTGAGCA 304
DB 25183 CTGGAGGCAAGAGTTGCAAGTGAAGTGAATGTCCTGACTCCAGCCTGGGCAACA 25242
QY 305 GAGTGAGACCKCGTCTCAAAAACAACAACAAAAA 347
DB 25243 GTGTGAGACTCTCATCTCAAAAAGAAATAAATAAATAAATAAATAA 25285
```

```
RESULT 36
US-09-949-016-13852
; Sequence 13852, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; FILE REFERENCE: CL001307
; CURRENT APPLICATION NUMBER: US/09/949,016
; PRIOR FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 13852
; LENGTH: 86945
; TYPE: DNA
; ORGANISM: Human
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: (1)...(86945)
; OTHER INFORMATION: n = A,T,C or G
US-09-949-016-13852

Query Match 37.0%; Score 147.8; DB 3; Length 86945;
Best Local Similarity 73.5%; Pred. No. 2.9e-33;
Matches 208; Conservative 1; Mismatches 58; Indels 16; Gaps 1;

QY 81 ATGCTGTAATCCAGCACTTCGGAGGCCAAGTGGCGGATGACCTGAGTCAAGA 140
DB 25003 ATGCTGTAATCCAGCACTTCGGAGGCCAAGTGGCGGATGACCTGAGTCAAGA 25062
QY 141 TCGAGACCATCTGCGCCAACTGTGAACCCCGTCTTACTTAAATAACAAAAATAGC 200
DB 25063 TCGAGACCATCTGCGCCAACTGTGAACCCCGTCTTACTTAAATAACAAAAATAGC 25122
QY 201 TGGGATGTTGGCAACACCTGTAGTCCAGCTACTCAGAGCCGAGATTGCAGTAGC 260
DB 25123 TGGGATGTTGGCAACACCTGTAGTCCAGCTACTCAGAGCCGAGATTGCAGTAGC 25182
QY 261 TGAGATCCAGAGTGAAGCCGAATACAGAT-----CACAGAGTGAGCA 304
DB 25183 CTGGAGGCAAGAGTTGCAAGTGAAGTGAATGTCCTGACTCCAGCCTGGGCAACA 25242
QY 305 GAGTGAGACCKCGTCTCAAAAACAACAACAAAAA 347
DB 25243 GTGTGAGACTCTCATCTCAAAAAGAAATAAATAAATAAATAAATAA 25285

RESULT 37
US-09-949-016-13853
; Sequence 13853, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; FILE REFERENCE: CL001307
; CURRENT APPLICATION NUMBER: US/09/949,016
; PRIOR FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
```



```
US-09-949-016-13856
; Sequence 13856, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
; FILE REFERENCE: C1001307
; CURRENT APPLICATION NUMBER: US/09/949,016
; CURRENT FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 13856
; LENGTH: 86945
; TYPE: DNA
; ORGANISM: Human
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: (1)...(86945)
; OTHER INFORMATION: n = A,T,C or G
US-09-949-016-13856

Query Match
Best Local Similarity 37.0%; Score 147.8; DB 3; Length 86945;
Best Local Similarity 73.5%; Pred. No. 2.9e-33;
Matches 208; Conservative 1; Mismatches 58; Indels 16; Gaps 1;

QY 81 ATGCTGTAATCCCGACACTTCGGAGGCGCAAGTGGGCGGATCACTGAGTCAAGAGA 140
DB 25003 ATGCTGTAATCCCGACACTTCGGAGGCGCAAGTGGGCGGATCACTGAGTCAAGAGA 140
QY 141 TCGAGACCATCTGGCCCAACATGTGTAAACCCCGTCTTACTTAAATAATCAAAAAATATGAC 200
DB 25063 TTGAGACCAAGCTGGCCCAACATGTGTAAACCCCGTCTTACTTAAATAATCAAAAAATATGAGC 25122
QY 201 TGGGCGATGTGGGCGACACACCTGTAGTCCAGCTACTCAGAGGCGGAGATTGCACTGAGC 260
DB 25123 TGGGCGATGTGGGCGACACACCTGTAGTCCAGCTACTCAGAGGCGGAGATTGCACTGAGC 25182
QY 261 TGAGATCGCAGAGTGAGCGGCAATCAAGAT-----CACAGAGTGAGCA 304
DB 25183 CTGGAGGCGCAGAGTGAGCGGCAATCAAGAT-----CACAGAGTGAGCA 25242
QY 305 GAGTGAGACKCGCTCTCAAAAACAACAACAAAAAACA 347
DB 25243 GTGTGAGACTCATCTCAAAAAGAATAAATAAATAATA 25285

RESULT 41
US-09-949-016-13857
; Sequence 13857, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
; FILE REFERENCE: C1001307
; CURRENT APPLICATION NUMBER: US/09/949,016
; CURRENT FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 13857
```

```
LENGTH: 86945
; TYPE: DNA
; ORGANISM: Human
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: (1)...(86945)
; OTHER INFORMATION: n = A,T,C or G
US-09-949-016-13857

Query Match
Best Local Similarity 37.0%; Score 147.8; DB 3; Length 86945;
Best Local Similarity 73.5%; Pred. No. 2.9e-33;
Matches 208; Conservative 1; Mismatches 58; Indels 16; Gaps 1;

QY 81 ATGCTGTAATCCCGACACTTCGGAGGCGCAAGTGGGCGGATCACTGAGTCAAGAGA 140
DB 25003 ATGCTGTAATCCCGACACTTCGGAGGCGCAAGTGGGCGGATCACTGAGTCAAGAGA 140
QY 141 TCGAGACCATCTGGCCCAACATGTGTAAACCCCGTCTTACTTAAATAATCAAAAAATATGAC 200
DB 25063 TTGAGACCAAGCTGGCCCAACATGTGTAAACCCCGTCTTACTTAAATAATCAAAAAATATGAGC 25122
QY 201 TGGGCGATGTGGGCGACACACCTGTAGTCCAGCTACTCAGAGGCGGAGATTGCACTGAGC 260
DB 25123 TGGGCGATGTGGGCGACACACCTGTAGTCCAGCTACTCAGAGGCGGAGATTGCACTGAGC 25182
QY 261 TGAGATCGCAGAGTGAGCGGCAATCAAGAT-----CACAGAGTGAGCA 304
DB 25183 CTGGAGGCGCAGAGTGAGCGGCAATCAAGAT-----CACAGAGTGAGCA 25242
QY 305 GAGTGAGACKCGCTCTCAAAAACAACAACAAAAAACA 347
DB 25243 GTGTGAGACTCATCTCAAAAAGAATAAATAAATAATA 25285

RESULT 42
US-09-949-016-13858
; Sequence 13858, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
; FILE REFERENCE: C1001307
; CURRENT APPLICATION NUMBER: US/09/949,016
; CURRENT FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 13858
; LENGTH: 86945
; TYPE: DNA
; ORGANISM: Human
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: (1)...(86945)
; OTHER INFORMATION: n = A,T,C or G
US-09-949-016-13858

Query Match
Best Local Similarity 37.0%; Score 147.8; DB 3; Length 86945;
Matches 208; Conservative 1; Mismatches 58; Indels 16; Gaps 1;

QY 81 ATGCTGTAATCCCGACACTTCGGAGGCGCAAGTGGGCGGATCACTGAGTCAAGAGA 140
DB 25003 ATGCTGTAATCCCGACACTTCGGAGGCGCAAGTGGGCGGATCACTGAGTCAAGAGA 140
QY 141 TCGAGACCATCTGGCCCAACATGTGTAAACCCCGTCTTACTTAAATAATCAAAAAATATGAC 200
DB 25063 TTGAGACCAAGCTGGCCCAACATGTGTAAACCCCGTCTTACTTAAATAATCAAAAAATATGAGC 25122
QY 201 TGGGCGATGTGGGCGACACACCTGTAGTCCAGCTACTCAGAGGCGGAGATTGCACTGAGC 260
DB 25123 TGGGCGATGTGGGCGACACACCTGTAGTCCAGCTACTCAGAGGCGGAGATTGCACTGAGC 25182
QY 261 TGAGATCGCAGAGTGAGCGGCAATCAAGAT-----CACAGAGTGAGCA 304
DB 25183 CTGGAGGCGCAGAGTGAGCGGCAATCAAGAT-----CACAGAGTGAGCA 25242
QY 305 GAGTGAGACKCGCTCTCAAAAACAACAACAAAAAACA 347
DB 25243 GTGTGAGACTCATCTCAAAAAGAATAAATAAATAATA 25285
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D8		25063	TTGAGACACAGCTGGCCAAATGGTAAAAACCCTGTCTACTAAAATACAAAAATTGAGC	25122
Q7		201	TGGGCATCGTGTCACAACAACCTGTATGTGCCAGCTACTCAGAGCCGGAGATTGCATGAGC	260
D8		25123	TGGCCGCTGGTGCTGTGTCACCTATATCTCTGCTACTCAGAGGCTGAGGTTGCATGTAAC	25182
Q7		261	TGAATGTCAGAGTAGAGCCGAATTCACAGAT-----CACAGGTAGCA	304
D8		25183	CTGGGAGGACAGAGGTTGCAGCTGAGCTGAGATTGTGCCATTGTACTCCAGCCTGGGCACA	25242
Q7		305	GAGTGAGACKCCGCTCAAAAACAACAACAAAAA :	347
D8		25243	GTGTGAGACTTCATCTCAAAAAGATATTAATTAATTAATTA	25285

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RESULT 43
US-09-949-016-14022
: Sequence 14022, Application US/09949016
: Patent No. 6812339
: GENERAL INFORMATION:
: APPLICANT: VENTER, J. Craig et al.
: TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
: TITLE OF INVENTION: WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
: FILE REFERENCE: CLO01107
: CURRENT APPLICATION NUMBER: US/09/949, 016
: CURRENT FILING DATE: 2000-04-14
: PRIOR APPLICATION NUMBER: 60/241,755
: PRIOR FILING DATE: 2000-10-20
: PRIOR APPLICATION NUMBER: 60/237,768
: PRIOR FILING DATE: 2000-10-03
: PRIOR APPLICATION NUMBER: 60/231,498
: PRIOR FILING DATE: 2000-09-08
: NUMBER OF SEQ ID NOS: 207012
: SOFTWARE: FASTSEQ for Windows Version 4.0
: SEQ ID NO 14022.
: LENGTH: 9798
: TYPE: DNA
: ORGANISM: Human
: US-09-949-016-14022

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Best Local Similarity	75.4%	Pred. No. 1.4e-33		
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			Gaps	1

QY	84	CCTGTAATCCCA	CGACTTCGGGAGGCCAAAGTGCGCGATCACTGAGGTCAAAGATCG	143
DB	964	CCTTAATCCCGG	CACTTGGGAAACCAAGGACGAGGTGGCTTAGGCCACGAGATTCA	1023
QY	144	AGAATCATCTG	AGCCCAATGGTGAACCCCGTCTTACTTAATAAATCAAAAAATTAAGCTGG	203
DB	1024	AGACCAAGCTT	GGCCCAATGGTGAACCCCGTCTTACTTAATAAATCAAAAAATTAAGCCGG	1083
QY	204	GCATGGTGGCA	CACACTCTGTAGTCCAGCTACTCAAGACCGGAGATTGCAATGAGCTGA	263
DB	1084	ACATGGTGGCA	AGTGCCTGTATATCCAGAGTACTCAAGAGTGGAGGTTGCAATGAGCGGA	1143
QY	264	GATGGCAGAGT	GAGCCGAAATACACAGATACACAGAGTGAAGACAGATGACCKCCGTCTCAA	323
DB	1144	GATCAACAGCC	CTCGCACTCCAGCTGGTGA-----CAAGCGAGACTCATCTCAA	1194
QY	324	AACGACACAC	AAAAAACAACAAAAA	347
DB	1195	AACGAAAAA	CAAAAAACAGACAA	1218

RESULT 44
US-09-949-016-12819
Sequence 12819, Application US/09949016
Patent No. 6812339
GENERAL INFORMATION:
APPLICANT: VENTER, J. Craig et al.
TITLE OF INVENTION: WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF

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: FILE REFERENCE: CL001307
: CURRENT APPLICATION NUMBER: US/09/949,016
: CURRENT FILING DATE: 2000-04-14
: PRIOR APPLICATION NUMBER: 60/241,755
: PRIOR FILING DATE: 2000-10-20
: PRIOR APPLICATION NUMBER: 60/237,768
: PRIOR FILING DATE: 2000-10-03
: PRIOR APPLICATION NUMBER: 60/231,498
: PRIOR FILING DATE: 2000-09-08
: NUMBER OF SEQ ID NOS: 207012
: SOFTWARE: FASTSQ for Windows Version 4.0
: SEQ ID NO 12819
: LENGTH: 9801
: TYPE: DNA
: ORGANISM: Human
: US-09-949-016-12819

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	Best Local Similarity	75.4%;	Pred. No. 1.4e-33;		
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Qy	144	AGACATCTCTGGCCCAATGATGTGAAGCCCGCTTTACTATAAAATATACAAAAAATATAGCTGG	203		
Db	1024	AGACCAACCTGGGCCCATGTGTGAAGCCCGCTCTACTATAAAATATACAAAAAATATAGCCGG	1081		
Qy	204	GCATGTGGGCACACACCTGTATGTCCAGCTACTCAGAGAGCCGGAGATTGCATGTAGCTGA	263		
Db	1084	ACATGTGTGGCACCTGTCTGTATATCCACAGTACTCAAGAGGTGGAGGTTGCAGGTGACCGGA	1144		
Qy	264	GATGCGAGAGTGTAGCCGAAATTCACAGATTCACAGATGTAGCAGAGTGTAGACCCGTCCTCA	323		
Db	1144	GATCAGACGCCCTGTGACTTCCAGCGCTGGGTGA-----CAGAGGAGACATTCATCTCA	1199		
Qy	324	AAACACACACAAAAAACAAAAA	347		
Db	1195	AACGAAAAACAAAAACAGACCA	1218		

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RESULT 45
US-09-936-271C-56
: Sequence 56 Application US/09936271C
: Patent No. 7022497
: GENERAL INFORMATION:
: APPLICANT: Youself, George M.
: APPLICANT: Diamandis, Eleftherios
: TITLE OF INVENTION: No. 7022497el Human Kallikrein-Like Genes
: FILE REFERENCE: WTSJUSA
: CURRENT APPLICATION NUMBER: US/09/936,271C
: CURRENT FILING DATE: 2001-09-10
: PRIOR APPLICATION NUMBER: PCT/CA00/00258
: PRIOR FILING DATE: 2000-03-09
: PRIOR APPLICATION NUMBER: US 60/124,260
: PRIOR FILING DATE: 1999-03-11
: PRIOR APPLICATION NUMBER: US 60/127,386
: PRIOR FILING DATE: 1999-04-01
: PRIOR APPLICATION NUMBER: US 60/144,919
: PRIOR FILING DATE: 1999-07-21
: NUMBER OF SEQ ID NOS: 97
: SOFTWARE: PatentIn version 3.2
: SEQ ID NO 56
: LENGTH: 11820
: TYPE: DNA
: ORGANISM: Homo sapiens
: US-09-936-271C-56

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Query Match	36.9%;	Score 147.6;	DB 5;	Length 11820;	.
Best Local Similarity	75.4%;	Pred. No. 1.5e-33;			
Matches 199; Conservative	1;	Mismatches 55;	Indels 9;	Gaps 1;	

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GenCore version 5.1.9
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Perfect score: 399.6

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Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

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4	397.2	99.4	75729	US-10-741-601-13958	Sequence 13958, A
5	199	49.8	201	US-10-741-601-13958	Sequence 13958, A
6	199	49.8	201	US-10-741-601-14097	Sequence 14097, A
7	199	49.8	201	US-10-741-601-14097	Sequence 14097, A
8	199	49.8	201	US-10-741-600-35944	Sequence 35944, A
9	197.2	49.3	201	US-10-741-600-35121	Sequence 35121, A
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11	197.2	49.3	201	US-10-741-601-14060	Sequence 14060, A
12	197.2	49.3	201	US-10-741-600-36084	Sequence 36084, A
13	185	46.3	201	US-10-741-601-13992	Sequence 13992, A
14	185	46.3	201	US-10-741-601-14059	Sequence 14059, A
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16	185	46.3	201	US-10-741-600-36083	Sequence 36083, Ap
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19	158.4	39.6	96593	US-09-997-722-16	Sequence 16, App
20	158.2	39.6	997	US-10-301-480-603905	Sequence 603905, A
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27	154.8	38.7	614	US-09-925-065A-532789	Sequence 532789, A
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29	153.6	38.4	977	US-10-301-480-609460	Sequence 609460, A
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37	152	38.0	181343	US-10-756-149-2215	Sequence 2215, Ap
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52	150.4	37.6	17538	US-10-810-653-9	Sequence 9, App
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82	149	37.3	8205	US-10-227-646-776	Sequence 276, App
83	149	37.3	89900	US-10-722-839-3	Sequence 3, App
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86	148.8	37.2	44075	US-09-997-722-10	Sequence 4075, App
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C 102	147.6	36.9	983	12	US-10-301-480-538078	Sequence 538078,
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C 106	147.6	36.9	180531	16	US-11-114-798-57	Sequence 57, Appl
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C 112	147.2	36.8	599001	9	US-10-317-869A-4	Sequence 4, Appl
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C 114	147.2	36.8	783062	10	US-10-461-862-166	Sequence 166, App
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ALIGNMENTS

RESULT 1
US-10-741-601-5648/c
Sequence 5648, Application US/10741601
Publication No. US20040166519A1
GENERAL INFORMATION:
APPLICANT: CARGILL, Michele et al.
TITLE OF INVENTION: GENETIC POLYMORPHISMS ASSOCIATED WITH
STENOSIS, METHODS OF DETECTION AND USES THEREOF

FILE REFERENCE: CL001500
CURRENT APPLICATION NUMBER: US/10/741,601
CURRENT FILING DATE: 2003-12-22
NUMBER OF SEQ ID NOS: 26415
SOFTWARE: FASTSEQ for Windows Version 4.0
SEQ ID NO 5648
LENGTH: 36805
TYPE: DNA
ORGANISM: Homo sapiens
FEATURE:
NAME/KEY: misc_feature
LOCATION: (1)...(36805)
OTHER INFORMATION: n = A,T,C or G, or insertion/deletion polymorphism (see Tables 1-
US-10-741-601-5648

Query Match 99.4%; Score 397.2; DB 8; Length 36805;
Best Local Similarity 99.2%; Pred. No. 2.2e-11;
Matches 397; Conservative 2; Mismatches 1; Indels 0; Gaps 0;

1 CCAAGTCTGACGCTGATGCTGGGCGCATGGGAAACCAATATTATTAAGACATTGTCAGG 60
DB CCAAGTCTGACGCTGATGCTGGGCGCATGGGAAACCAATATTATTAAGACATTGTCAGG 27741
61 CCAAGCATGACACTGCTGTAATGCTGTAATCCAGCACTTCGGAGGCCAAGGTGGCG 120
DB CCAAGCATGACACTGCTGTAATGCTGTAATCCAGCACTTCGGAGGCCAAGGTGGCG 27681
121 GATACCTGAGGTGAAGATCCAGACCATCTCTGGCCAACTGGTGAACCCCTCTTTA 180
DB GATACCTGAGGTGAAGATCCAGACCATCTCTGGCCAACTGGTGAACCCCTCTTTA 27621
27680 GATACCTGAGGTGAAGATCCAGACCATCTCTGGCCAACTGGTGAACCCCTCTTTA 27621
181 CTAATAATACAAAAATAGCTGGGCGATGTCGACACACCTGTATCCAGCTACTGAGG 240
DB CTAATAATACAAAAATAGCTGGGCGATGTCGACACACCTGTATCCAGCTACTGAGG 27620
27620 CTAATAATACAAAAATAGCTGGGCGATGTCGACACACCTGTATCCAGCTACTGAGG 27561
241 AGCGGAGATTGCAATGCTGATGATCGAGATGAGCCGAATCAAGATCAGAGTGG 300
DB AGCGGAGATTGCAATGCTGATGATCGAGATGAGCCGAATCAAGATCAGAGTGG 27560
27560 AGCGGAGATTGCAATGCTGATGATCGAGATGAGCCGAATCAAGATCAGAGTGG 27501
301 AGCAGATGAGACCCCTCTTAAACCAACCAACCAACCAACCAACCAACCAACCAACCA 360
DB AGCAGATGAGACCCCTCTTAAACCAACCAACCAACCAACCAACCAACCAACCAACCA 27500
27500 AGCAGATGAGACCCCTCTTAAACCAACCAACCAACCAACCAACCAACCAACCAACCA 27441
361 TCCATCTGCGGTCCCACTATTGACGAGACCAAAAG 400
DB TCCATCTGCGGTCCCACTATTGACGAGACCAAAAG 27440
27440 TCCATCTGCGGTCCCACTATTGACGAGACCAAAAG 27401

RESULT 2
US-10-741-600-17657/c
Sequence 17657, Application US/10741600
Publication No. US20050026169A1
GENERAL INFORMATION:
APPLICANT: CARGILL, Michele et al.
TITLE OF INVENTION: GENETIC POLYMORPHISMS ASSOCIATED WITH
MYOCARDIAL INFARCTION, METHODS OF DETECTION AND USES THEREOF
FILE REFERENCE: CL001499
CURRENT APPLICATION NUMBER: US/10/741,600
CURRENT FILING DATE: 2003-12-22
NUMBER OF SEQ ID NOS: 73997
SOFTWARE: FASTSEQ for Windows Version 4.0
SEQ ID NO 17657
LENGTH: 36805
TYPE: DNA
ORGANISM: Homo sapiens
FEATURE:
NAME/KEY: misc_feature
LOCATION: (1)...(36805)
OTHER INFORMATION: n = A,T,C or G, or insertion/deletion polymorphism (see Tables 1-
US-10-741-600-17657

Query Match 99.4%; Score 397.2; DB 9; Length 36805;
Best Local Similarity 99.2%; Pred. No. 2.2e-11;

Matches 397; Conservative 2; Mismatches 1; Indels 0; Gaps 0;

QY 1 CCAGGTAATCAGGCACTGCTGGCCATGCGAACCCTAATATTATTAAGACATTGTCCAG 60
 DB 27800 CCAGGTAATCAGGCACTGCTGGCCATGCGAACCCTAATATTATTAAGACATTGTCCAG 27741

QY 61 CCAGGCAATGACACTGGCTGTAATGCTGTATCCAGCACTTGGGAGGCCAAGTGGGCG 120
 DB 27740 CCAGGCAATGACACTGGCTGTAATGCTGTATCCAGCACTTGGGAGGCCAAGTGGGCG 27681

QY 121 GATCACTGAGGTCAAGAGATGAGACCATCTGGCCCAATGATGTGAACCCCGTCTTTA 180
 DB 27680 GATCACTGAGGTCAAGAGATGAGACCATCTGGCCCAATGATGTGAACCCCGTCTTTA 27621

QY 181 CTAATAATACAAAAATAGCTGGGCAATGCTGGCAACACCTGTATCCCACTACTCAG 240
 DB 27620 CTAATAATACAAAAATAGCTGGGCAATGCTGGCAACACCTGTATCCCACTACTCAG 27561

QY 241 AGCCGAGATTGCACTGAGCTGAGATCGCAGAGTGGAGCCGAATGACAGATCAGAGAGTG 300
 DB 27560 AGCCGAGATTGCACTGAGCTGAGATCGCAGAGTGGAGCCGAATGACAGATCAGAGAGTG 27501

QY 301 AGCAGAGTGAAGACCCGCTCTCAAAAAACAACAAAAACAAACCATTAAGACATTG 360
 DB 27500 AGCAGAGTGAAGACCCGCTCTCAAAAAACAACAAAAACAAACCATTAAGACATTG 27441

QY 361 TCCATCTGCGGTTCCCACTATTGTCAGAGACCAAAAG 400
 DB 27440 TCCATCTGCGGTTCCCACTATTGTCAGAGACCAAAAG 27401

RESULT 3
 US-10-741-601-5649/c
 ; Sequence 5649, Application US/10741601
 ; Publication No. US20040166519A1
 ; GENERAL INFORMATION:
 ; APPLICANT: CARGILL, Michele et al.
 ; TITLE OF INVENTION: GENETIC POLYMORPHISMS ASSOCIATED WITH
 ; TITLE OF INVENTION: STENOSIS, METHODS OF DETECTION AND USES THEREOF
 ; FILE REFERENCE: CLO01500
 ; CURRENT APPLICATION NUMBER: US/10/741,601
 ; CURRENT FILING DATE: 2003-12-22
 ; NUMBER OF SEQ ID NOS: 26415
 ; SOFTWARE: FASTSEQ for Windows Version 4.0
 ; SEQ ID NO: 5649
 ; LENGTH: 75729
 ; TYPE: DNA
 ; ORGANISM: Homo sapiens
 ; FEATURE:
 ; NAME/KEY: misc_feature
 ; LOCATION: (1)...(75729)
 ; OTHER INFORMATION: n = A,T,C or G, or insertion/deletion polymorphism (see Tables 1-
 US-10-741-601-5649

Query Match 99.4%; Score 397.2; DB 8; Length 75729;
 Best Local Similarity 99.2%; Pred. No. 3e-111; Indels 0; Gaps 0;
 Matches 397; Conservative 2; Mismatches 1; Indels 0; Gaps 0;

QY 1 CCAGGTAATCAGGCACTGCTGGCCATGCGAACCCTAATATTATTAAGACATTGTCCAG 60
 DB 59242 CCAGGTAATCAGGCACTGCTGGCCATGCGAACCCTAATATTATTAAGACATTGTCCAG 59183

QY 61 CCAGGCAATGACACTGGCTGTAATGCTGTATCCAGCACTTGGGAGGCCAAGTGGGCG 120
 DB 59182 CCAGGCAATGACACTGGCTGTAATGCTGTATCCAGCACTTGGGAGGCCAAGTGGGCG 59123

QY 121 GATCACTGAGGTCAAGAGATGAGACCATCTGGCCCAATGATGTGAACCCCGTCTTTA 180
 DB 59122 GATCACTGAGGTCAAGAGATGAGACCATCTGGCCCAATGATGTGAACCCCGTCTTTA 59063

QY 181 CTAATAATACAAAAATAGCTGGGCAATGCTGGCAACACCTGTATCCCACTACTCAG 240
 DB 59062 CTAATAATACAAAAATAGCTGGGCAATGCTGGCAACACCTGTATCCCACTACTCAG 59003

QY 241 AGCCGAGATTGCACTGAGCTGAGATCGCAGAGTGGAGCCGAATGACAGATCAGAGAGTG 300
 DB 59002 AGCCGAGATTGCACTGAGCTGAGATCGCAGAGTGGAGCCGAATGACAGATCAGAGAGTG 58943

QY 301 AGCAGAGTGAAGACCCGCTCTCAAAAAACAACAAAAACAAACCATTAAGACATTG 360
 DB 58942 AGCAGAGTGAAGACCCGCTCTCAAAAAACAACAAAAACAAACCATTAAGACATTG 58883

QY 361 TCCATCTGCGGTTCCCACTATTGTCAGAGACCAAAAG 400
 DB 58882 TCCATCTGCGGTTCCCACTATTGTCAGAGACCAAAAG 58843

RESULT 4
 US-10-741-600-17658/c
 ; Sequence 17658, Application US/10741600
 ; Publication No. US2005002619A1
 ; GENERAL INFORMATION:
 ; APPLICANT: CARGILL, Michele et al.
 ; TITLE OF INVENTION: GENETIC POLYMORPHISMS ASSOCIATED WITH
 ; TITLE OF INVENTION: MYOCARDIAL INFARCTION, METHODS OF DETECTION AND USES THEREOF
 ; FILE REFERENCE: CLO01499
 ; CURRENT APPLICATION NUMBER: US/10/741,600
 ; CURRENT FILING DATE: 2003-12-22
 ; NUMBER OF SEQ ID NOS: 73397
 ; SOFTWARE: FASTSEQ for Windows Version 4.0
 ; SEQ ID NO: 17658
 ; LENGTH: 75729
 ; TYPE: DNA
 ; ORGANISM: Homo sapiens
 ; FEATURE:
 ; NAME/KEY: misc_feature
 ; LOCATION: (1)...(75729)
 ; OTHER INFORMATION: n = A,T,C or G, or insertion/deletion polymorphism (see Tables 1-
 US-10-741-600-17658

Query Match 99.4%; Score 397.2; DB 9; Length 75729;
 Best Local Similarity 99.2%; Pred. No. 3e-111; Indels 0; Gaps 0;
 Matches 397; Conservative 2; Mismatches 1; Indels 0; Gaps 0;

QY 1 CCAGGTAATCAGGCACTGCTGGCCATGCGAACCCTAATATTATTAAGACATTGTCCAG 60
 DB 59242 CCAGGTAATCAGGCACTGCTGGCCATGCGAACCCTAATATTATTAAGACATTGTCCAG 59183

QY 61 CCAGGCAATGACACTGGCTGTAATGCTGTATCCAGCACTTGGGAGGCCAAGTGGGCG 120
 DB 59182 CCAGGCAATGACACTGGCTGTAATGCTGTATCCAGCACTTGGGAGGCCAAGTGGGCG 59123

QY 121 GATCACTGAGGTCAAGAGATGAGACCATCTGGCCCAATGATGTGAACCCCGTCTTTA 180
 DB 59122 GATCACTGAGGTCAAGAGATGAGACCATCTGGCCCAATGATGTGAACCCCGTCTTTA 59063

QY 181 CTAATAATACAAAAATAGCTGGGCAATGCTGGCAACACCTGTATCCCACTACTCAG 240
 DB 59062 CTAATAATACAAAAATAGCTGGGCAATGCTGGCAACACCTGTATCCCACTACTCAG 59003

QY 241 AGCCGAGATTGCACTGAGCTGAGATCGCAGAGTGGAGCCGAATGACAGATCAGAGAGTG 300
 DB 59002 AGCCGAGATTGCACTGAGCTGAGATCGCAGAGTGGAGCCGAATGACAGATCAGAGAGTG 58943

QY 301 AGCAGAGTGAAGACCCGCTCTCAAAAAACAACAAAAACAAACCATTAAGACATTG 360
 DB 58942 AGCAGAGTGAAGACCCGCTCTCAAAAAACAACAAAAACAAACCATTAAGACATTG 58883

QY 361 TCCATCTGCGGTTCCCACTATTGTCAGAGACCAAAAG 400
 DB 58882 TCCATCTGCGGTTCCCACTATTGTCAGAGACCAAAAG 58843

RESULT 5
 US-10-741-601-13924/c
 ; Sequence 13924, Application US/10741601

```
Publication No. US20040166519A1
; GENERAL INFORMATION:
; APPLICANT: CARGILL, Michele et al.
; TITLE OF INVENTION: GENETIC POLYMORPHISMS ASSOCIATED WITH
; FILE REFERENCE: CLO01500
; CURRENT APPLICATION NUMBER: US/10/741,601
; CURRENT FILING DATE: 2003-12-22
; NUMBER OF SEQ ID NOS: 26415
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 13924
; LENGTH: 201
; TYPE: DNA
; ORGANISM: Homo sapiens
US-10-741-601-13924

Query Match          49.8%; Score 199; DB 8; Length 201;
Best Local Similarity 99.0%; Pred. No. 5.9e-51;
Matches 199; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

QY 27 ATGGAAACCCAAATTTAATAAGACATTGTCAAGCCAGGCAATGACACTGGCTGAATGCTT 86
DB 201 ATGGAAACCCAAATTTAATAAGACATTGTCAAGCCAGGCAATGACACTGGCTGAATGCTT 142
QY 87 GTAATCCAGCACTTCGGAGAGCCCAAGGTGGCGGATCACTGAGGTCAAGAGATCGAGA 146
DB 141 GTAATCCAGCACTTCGGAGAGCCCAAGGTGGCGGATCACTGAGGTCAAGAGATCGAGA 82
QY 147 CCATCTGGCCCAACATGCTGAAACCCCGCTTTACTTAATAAAATACAAAAATAGCTGGGCA 206
DB 81 CCATCTGGCCCAACATGCTGAAACCCCGCTTTACTTAATAAAATACAAAAATAGCTGGGCA 22
QY 207 TGGTGGCACACACTGTAGTC 227
DB 21 TGGTGGCACACACTGTAGTC 1

RESULT 6
US-10-741-601-14097/c
; Sequence 14097, Application US/10741601
; Publication No. US20040166519A1
; GENERAL INFORMATION:
; APPLICANT: CARGILL, Michele et al.
; TITLE OF INVENTION: GENETIC POLYMORPHISMS ASSOCIATED WITH
; FILE REFERENCE: CLO01500
; CURRENT APPLICATION NUMBER: US/10/741,601
; CURRENT FILING DATE: 2003-12-22
; NUMBER OF SEQ ID NOS: 26415
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 14097
; LENGTH: 201
; TYPE: DNA
; ORGANISM: Homo sapiens
US-10-741-601-14097

Query Match          49.8%; Score 199; DB 8; Length 201;
Best Local Similarity 99.0%; Pred. No. 5.9e-51;
Matches 199; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

QY 27 ATGGAAACCCAAATTTAATAAGACATTGTCAAGCCAGGCAATGACACTGGCTGAATGCTT 86
DB 201 ATGGAAACCCAAATTTAATAAGACATTGTCAAGCCAGGCAATGACACTGGCTGAATGCTT 142
QY 87 GTAATCCAGCACTTCGGAGAGCCCAAGGTGGCGGATCACTGAGGTCAAGAGATCGAGA 146
DB 141 GTAATCCAGCACTTCGGAGAGCCCAAGGTGGCGGATCACTGAGGTCAAGAGATCGAGA 82
QY 147 CCATCTGGCCCAACATGCTGAAACCCCGCTTTACTTAATAAAATACAAAAATAGCTGGGCA 206
DB 81 CCATCTGGCCCAACATGCTGAAACCCCGCTTTACTTAATAAAATACAAAAATAGCTGGGCA 22
QY 207 TGGTGGCACACACTGTAGTC 227
```

```
DB 21 TGGTGGCACACACTGTAGTC 1

RESULT 7
US-10-741-600-35944/c
; Sequence 35944, Application US/10741600
; Publication No. US20050026169A1
; GENERAL INFORMATION:
; APPLICANT: CARGILL, Michele et al.
; TITLE OF INVENTION: GENETIC POLYMORPHISMS ASSOCIATED WITH
; FILE REFERENCE: CLO01499
; CURRENT APPLICATION NUMBER: US/10/741,600
; CURRENT FILING DATE: 2003-12-22
; NUMBER OF SEQ ID NOS: 73997
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 35944
; LENGTH: 201
; TYPE: DNA
; ORGANISM: Homo sapiens
US-10-741-600-35944

Query Match          49.8%; Score 199; DB 9; Length 201;
Best Local Similarity 99.0%; Pred. No. 5.9e-51;
Matches 199; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

QY 27 ATGGAAACCCAAATTTAATAAGACATTGTCAAGCCAGGCAATGACACTGGCTGAATGCTT 86
DB 201 ATGGAAACCCAAATTTAATAAGACATTGTCAAGCCAGGCAATGACACTGGCTGAATGCTT 142
QY 87 GTAATCCAGCACTTCGGAGAGCCCAAGGTGGCGGATCACTGAGGTCAAGAGATCGAGA 146
DB 141 GTAATCCAGCACTTCGGAGAGCCCAAGGTGGCGGATCACTGAGGTCAAGAGATCGAGA 82
QY 147 CCATCTGGCCCAACATGCTGAAACCCCGCTTTACTTAATAAAATACAAAAATAGCTGGGCA 206
DB 81 CCATCTGGCCCAACATGCTGAAACCCCGCTTTACTTAATAAAATACAAAAATAGCTGGGCA 22
QY 207 TGGTGGCACACACTGTAGTC 227
DB 21 TGGTGGCACACACTGTAGTC 1

RESULT 8
US-10-741-600-36121/c
; Sequence 36121, Application US/10741600
; Publication No. US20050026169A1
; GENERAL INFORMATION:
; APPLICANT: CARGILL, Michele et al.
; TITLE OF INVENTION: MYOCARDIAL INFARCTION, METHODS OF DETECTION AND USES THEREOF
; FILE REFERENCE: CLO01499
; CURRENT APPLICATION NUMBER: US/10/741,600
; CURRENT FILING DATE: 2003-12-22
; NUMBER OF SEQ ID NOS: 73997
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 36121
; LENGTH: 201
; TYPE: DNA
; ORGANISM: Homo sapiens
US-10-741-600-36121

Query Match          49.8%; Score 199; DB 9; Length 201;
Best Local Similarity 99.0%; Pred. No. 5.9e-51;
Matches 199; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

QY 27 ATGGAAACCCAAATTTAATAAGACATTGTCAAGCCAGGCAATGACACTGGCTGAATGCTT 86
DB 201 ATGGAAACCCAAATTTAATAAGACATTGTCAAGCCAGGCAATGACACTGGCTGAATGCTT 142
QY 87 GTAATCCAGCACTTCGGAGAGCCCAAGGTGGCGGATCACTGAGGTCAAGAGATCGAGA 146
```

D5	141	GTATATCCACACTTTGGGAGGCCAAGTGGCGGATCACTGAAGTCAAGAGATCGAGA	82
OY	147	CCATCTCTGGCCAACTATGTAAACCCCGTCTTTACTPAAAAATPACAAAAATATGCTGGGCA	20
D6	81	CCATCTCTGGCCAACTATGTAAACCCCGTCTTTACTPAAAAATPACAAAAATATGCTGGGCA	22
OY	207	TGGTGGGACACACTGTATGTC	227
D6	21	TGGTGGGACACACTGTATGTC	1

RESULT 9
10-74

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US-10-741-601-13993/c
? Sequence 13993, Application US/10741601
? Publication No. US2004016551A1
? GENERAL INFORMATION:
? APPLICANT: CARGILL, Michele et al.
? TITLE OF INVENTION: GENETIC POLYMORPHISMS ASSOCIATED WITH
? TITLE OF INVENTION: STENOSIS, METHODS OF DETECTION AND USES THEREOF
? FILE REFERENCE: CLO01500
? CURRENT APPLICATION NUMBER: US/10/741,601
? CURRENT FILING DATE: 2003-12-22
? NUMBER OF SEQ ID NOS: 26415
? SOFTWARE: FastSeq for Windows Version 4.0
? SEQ ID NO 13993.
? LENGTH: 201
? TYPE: DNA
? ORGANISM: Homo sapiens
US-10-741-601-13993

```

Query Match	49.3%	Score 197.2;	DB 8;	Length 201;
Best Local Similarity	99.0%	Pred. No. 2.1e-50;		
Matches 196; Conservative	2;	Mismatches 0;	Indels 0;	Gaps 0

[illegible]

RESULT 10

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US-10-741-601-14060/c
Sequence 14060, Application US/10741601
Publication No. US20040166519A1
GENERAL INFORMATION:
APPLICANT: CARGILL, Michèle et al.
TITLE OF INVENTION: GENETIC POLYMORPHISMS ASSOCIATED WITH
FILE REFERENCE: CL001500
CURRENT APPLICATION NUMBER: US/10/741,601
CURRENT FILING DATE: 2003-12-22
NUMBER OF SEQ ID NOS: 26415
SOFTWARE: FastSeq for Windows Version 4.0
SEQ ID NO 14060
LENGTH: 201
TYPE: DNA
ORGANISM: Homo sapiens
US-10-741-601-14060

```

Query Match	49.3%	Score 197.2;	DB 8;	Length 201;
Best Local Similarity	99.0%	Pred. No. 2.1e-50;		
Matches 196; Conservative	2;	Mismatches 0;	Indels 0;	Gaps 0;

Qy	203	GGGATGGTGGCA	CACACCTGTAGTCCAGACTCA	CAGAGGCCGAGATTCCAGTGAAGCTG	262
Db	201	GGCATGGTGGCA	CACACTGTAGTCCAGACTCA	CTCAGAGGCCGAGATTCCAGTGAAGCTG	142
Qy	263	AGATCCGAGGTGAG	CGCGAAATCACAGATCA	CAGAGTGAGCAGAGTGAAGCCKCGTCTCA	322
Db	141	AGATCCGAGGTGAG	CGCGAAATCACAGATCA	CAGAGTGAGGAGAGTGAAGTCCGTTCTCA	82
Qy	323	AAAAACAACA	CAAAAAACCAAAAAACCATTAAGACATTTGTCATTTGGGGTTTCCCAACTA		382
Db	81	AAAAACAACA	CAAAAAACCAAAAAACCATTAAGACATTTGTCATTTGGGGTTTCCCAACTA		22
Qy	383	TTGCAGAGAGCA	CAAAAAG	400	
Db	21	TTGCAGAGAGCA	CAAAAAG	4	

RESULT 11

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US-10-741-600-36017/C
; Sequence 36017, Application US/10741600
; Publication No. US20050026169A1
; GENERAL INFORMATION:
; APPLICANT: CARGILL, Michele et al.
; TITLE OF INVENTION: GENETIC POLYMORPHISMS ASSOCIATED WITH
; TITLE OF INVENTION: MYOCARDIAL INFARCTION, METHODS OF DETECTION AND USES THEREOF
; FILE REFERENCE: CLO01499
; CURRENT APPLICATION NUMBER: US/10/741,600
; CURRENT FILING DATE: 2003-12-22
; NUMBER OF SEQ ID NOS: 73997
; SOFTWARE: FastSeq for Windows Version 4.0
1. SEQ ID NO 36017
; LENGTH: 201
; TYPE: DNA
; ORGANISM: Homo sapiens
US-10-741-600-36017

```

Query Match	49.3%	Score 197.2;	DB 9;	Length 201;
Best Local Similarity	99.0%	Pred. No. 2.1e-50;		
Matches 196; Conservative	2;	Mismatches 0;	Indels 0;	Gaps 0;

Qy	203	GGCATTGGTGGCACACCTGTAGTCCAGCTACTCAGAACCCGAGATTGCATGTAGCTG	262
Db	201	GGCATTGGTGGCACACCTGTAGTCCAGCTACTCAGAACCCGAGATTGCATGTAGCTG	142
Qy	263	AGATGCGAGAGTGAGCCGAAATCA	322
Db	141	AGATGCGAGAGTGAGCCGAAATCA	82
Qy	323	AAAAACAACAATAAAAAACAATAAACAATTGTTCATCTGCGGTTCCAGACTA	382
Db	81	AAAAACAACAATAAAAAACAATAAACAATTGTTCATCTGCGGTTCCAGACTA	22
Qy	383	TTTGCAGAGAGCCAAAAAG	400
Db	21	TTTGCAGAGAGCCAAAAAG	4

RESULT 12

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US-10-/741-600/6084/C
; Sequence 36084, Application US/10741600
; Publication No. US20050026169A1
; GENERAL INFORMATION:
; APPLICANT: CARGILL, Michele et al.
; TITLE OF INVENTION: GENETIC POLYMORPHISMS ASSOCIATED WITH
; TITLE OF INVENTION: MYOCARDIAL INFARCTION, METHODS OF DETECTION AND USES THEREOF
; FILE REFERENCE: CL001499
; CURRENT APPLICATION NUMBER: US/10/741,600
; CURRENT FILING DATE: 2003-12-22
; NUMBER OF SEQ. ID NOS: 73997
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 36084
; LENGTH: 201

```

```

; TYPE: DNA
; ORGANISM: Homo sapiens
US-10-741-600-36084

Query Match
Best Local Similarity 49.3%; Score 197.2; DB 9; Length 201;
Matches 196; Conservative 2; Mismatches 0; Indels 0; Gaps 0;

QY 203 GGCATGTTGGCAGACACCTGTAGTCCAGCTACTCAGAGCCGGAGATTGAGTGAAGCTG 262
DB 201 GGCATGTTGGCAGACACCTGTAGTCCAGCTACTCAGAGCCGGAGATTGAGTGAAGCTG 142
QY 263 AGATCGCAGAGTGAAGCCGAATTCACAGATCAGAGTGAAGAGTGAAGTGAAGTGAAGTGA 322
DB 141 AGATCGCAGAGTGAAGCCGAATTCACAGATCAGAGTGAAGAGTGAAGTGAAGTGAAGTGA 82
QY 323 AAAACAACAACAAAAAACAATTAAGATCATTGCTGCGGTTCCAGACTA 382
DB 81 AAAACAACAACAAAAAACAATTAAGATCATTGCTGCGGTTCCAGACTA 22
QY 383 TTGCAGGAGACCAAAAAG 400
DB 21 TTGCAGGAGACCAAAAAG 4

RESULT 13
US-10-741-601-13992/c
; Sequence 13992, Application US/10741601
; Publication No. US20040166519A1
; GENERAL INFORMATION:
; APPLICANT: CARGILL, Michele et al.
; TITLE OF INVENTION: GENETIC POLYMORPHISMS ASSOCIATED WITH
; FILE REFERENCE: CL001500
; CURRENT APPLICATION NUMBER: US/10/741,601
; CURRENT FILING DATE: 2003-12-22
; NUMBER OF SEQ ID NOS: 26415
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 13992
; LENGTH: 201
; TYPE: DNA
; ORGANISM: Homo sapiens
US-10-741-601-13992

Query Match
Best Local Similarity 46.3%; Score 185; DB 8; Length 201;
Matches 186; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 214 ACACACCTGTAGTCCAGCTACTCAGAGCCGGAGATTGAGTGAAGTGAAGTGAAGTGAAGTGA 273
DB 201 ACACACCTGTAGTCCAGCTACTCAGAGCCGGAGATTGAGTGAAGTGAAGTGAAGTGAAGTGA 142
QY 274 TGAGCCGAATTCACAGATCAGAGTGAAGAGTGAAGTGAAGTGAAGTGAAGTGAAGTGAAGTGA 333
DB 141 TGAGCCGAATTCACAGATCAGAGTGAAGAGTGAAGTGAAGTGAAGTGAAGTGAAGTGAAGTGA 82
QY 334 AAAAAACAAAAAACAATTAAGATCATTGCTGCGGTTCCAGACTAATTGCGAGAGAC 393
DB 81 AAAAAACAAAAAACAATTAAGATCATTGCTGCGGTTCCAGACTAATTGCGAGAGAC 22
QY 394 CAAAAAG 400
DB 21 CAAAAAG 15

RESULT 14
US-10-741-601-14059/c
; Sequence 14059, Application US/10741601
; Publication No. US20040166519A1
; GENERAL INFORMATION:
; APPLICANT: CARGILL, Michele et al.
; TITLE OF INVENTION: GENETIC POLYMORPHISMS ASSOCIATED WITH
; FILE REFERENCE: CL001500
; CURRENT APPLICATION NUMBER: US/10/741,601
; CURRENT FILING DATE: 2003-12-22
; NUMBER OF SEQ ID NOS: 26415
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 13992
; LENGTH: 201
; TYPE: DNA
; ORGANISM: Homo sapiens
US-10-741-601-13992

Query Match
Best Local Similarity 46.3%; Score 185; DB 8; Length 201;
Matches 186; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 214 ACACACCTGTAGTCCAGCTACTCAGAGCCGGAGATTGAGTGAAGTGAAGTGAAGTGAAGTGA 273
DB 201 ACACACCTGTAGTCCAGCTACTCAGAGCCGGAGATTGAGTGAAGTGAAGTGAAGTGAAGTGA 142
QY 274 TGAGCCGAATTCACAGATCAGAGTGAAGAGTGAAGTGAAGTGAAGTGAAGTGAAGTGAAGTGA 333
DB 141 TGAGCCGAATTCACAGATCAGAGTGAAGAGTGAAGTGAAGTGAAGTGAAGTGAAGTGAAGTGA 82
QY 334 AAAAAACAAAAAACAATTAAGATCATTGCTGCGGTTCCAGACTAATTGCGAGAGAC 393
DB 81 AAAAAACAAAAAACAATTAAGATCATTGCTGCGGTTCCAGACTAATTGCGAGAGAC 22
QY 394 CAAAAAG 400
DB 21 CAAAAAG 15

RESULT 15
US-10-741-600-36016/c
; Sequence 36016, Application US/10741600
; Publication No. US20050026169A1
; GENERAL INFORMATION:
; APPLICANT: CARGILL, Michele et al.
; TITLE OF INVENTION: MYOCARDIAL INFARCTION, METHODS OF DETECTION AND USES THEREOF
; FILE REFERENCE: CL001499
; CURRENT APPLICATION NUMBER: US/10/741,600
; CURRENT FILING DATE: 2003-12-22
; NUMBER OF SEQ ID NOS: 73997
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 36016
; LENGTH: 201
; TYPE: DNA
; ORGANISM: Homo sapiens
US-10-741-600-36016

Query Match
Best Local Similarity 46.3%; Score 185; DB 9; Length 201;
Matches 186; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 214 ACACACCTGTAGTCCAGCTACTCAGAGCCGGAGATTGAGTGAAGTGAAGTGAAGTGAAGTGA 273
DB 201 ACACACCTGTAGTCCAGCTACTCAGAGCCGGAGATTGAGTGAAGTGAAGTGAAGTGAAGTGA 142
QY 274 TGAGCCGAATTCACAGATCAGAGTGAAGAGTGAAGTGAAGTGAAGTGAAGTGAAGTGAAGTGA 333
DB 141 TGAGCCGAATTCACAGATCAGAGTGAAGAGTGAAGTGAAGTGAAGTGAAGTGAAGTGAAGTGA 82
QY 334 AAAAAACAAAAAACAATTAAGATCATTGCTGCGGTTCCAGACTAATTGCGAGAGAC 393
DB 81 AAAAAACAAAAAACAATTAAGATCATTGCTGCGGTTCCAGACTAATTGCGAGAGAC 22
QY 394 CAAAAAG 400
DB 21 CAAAAAG 15

RESULT 16
US-10-741-601-14059/c
; Sequence 14059, Application US/10741601
; Publication No. US20040166519A1
; GENERAL INFORMATION:
; APPLICANT: CARGILL, Michele et al.
; TITLE OF INVENTION: GENETIC POLYMORPHISMS ASSOCIATED WITH
; FILE REFERENCE: CL001500
; CURRENT APPLICATION NUMBER: US/10/741,601
; CURRENT FILING DATE: 2003-12-22
; NUMBER OF SEQ ID NOS: 26415
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 14059
; LENGTH: 201
; TYPE: DNA
; ORGANISM: Homo sapiens
US-10-741-601-14059

Query Match
Best Local Similarity 46.3%; Score 185; DB 8; Length 201;
Matches 186; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 214 ACACACCTGTAGTCCAGCTACTCAGAGCCGGAGATTGAGTGAAGTGAAGTGAAGTGAAGTGA 273
DB 201 ACACACCTGTAGTCCAGCTACTCAGAGCCGGAGATTGAGTGAAGTGAAGTGAAGTGAAGTGA 142
QY 274 TGAGCCGAATTCACAGATCAGAGTGAAGAGTGAAGTGAAGTGAAGTGAAGTGAAGTGAAGTGA 333
DB 141 TGAGCCGAATTCACAGATCAGAGTGAAGAGTGAAGTGAAGTGAAGTGAAGTGAAGTGAAGTGA 82
QY 334 AAAAAACAAAAAACAATTAAGATCATTGCTGCGGTTCCAGACTAATTGCGAGAGAC 393
DB 81 AAAAAACAAAAAACAATTAAGATCATTGCTGCGGTTCCAGACTAATTGCGAGAGAC 22
QY 394 CAAAAAG 400
DB 21 CAAAAAG 15
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US-10-741-600-36083/c
; Sequence 36083, Application US/10741600
; Publication No. US20050026169A1
; GENERAL INFORMATION:
; APPLICANT: CARILL, Michele et al.
; TITLE OF INVENTION: GENETIC POLYMORPHISMS ASSOCIATED WITH
; FILE OF INVENTION: MYOCARDIAL INFARCTION, METHODS OF DETECTION AND USES THEREOF
; FILE REFERENCE: CLO001499
; CURRENT APPLICATION NUMBER: US/10/741,600
; CURRENT FILING DATE: 2003-12-22
; NUMBER OF SEQ ID NOS: 73997
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 36083
; LENGTH: 201
; TYPE: DNA
; ORGANISM: Homo sapiens
US-10-741-600-36083

Query Match          46.3% Score 185; DB 9; Length 201;
Best Local Similarity 99.5% Pred. No. 1.3e-46;
Matches 186; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 214 ACACACTGTAGTCCAGCTACTCTCAGAGCCGAGATTGCACTGAGTCCAGAG 273
DB 201 ACACACTGTAGTCCAGCTACTCTCAGAGCCGAGATTGCACTGAGTCCAGAG 142
QY 274 TGAGCCGAATTCACGATTCACAGAGTGAAGAGTGAACCCGCTCTCAAAAACAAAC 333
DB 141 TGAGCCGAATTCACGATTCACAGAGTGAAGAGTGAACCCGCTCTCAAAAACAAAC 82
QY 334 AAAAAACAAAAACCATTAAGCATTCCTGCGGTTCCCACTATTCAGAGAGC 393
DB 81 AAAAAACAAAAACCATTAAGCATTCCTGCGGTTCCCACTATTCAGAGAGC 22
QY 394 CAAAAAG 400
DB 21 CAAAAAG 15

RESULT 17
US-09-764-891-6505
; Sequence 6505, Application US/09764891
; Publication No. US20030077808A1
; GENERAL INFORMATION:
; APPLICANT: Rosen et al.
; TITLE OF INVENTION: Nucleic Acids, Proteins, and Antibodies
; FILE REFERENCE: PC006
; CURRENT APPLICATION NUMBER: US/09/764,891
; CURRENT FILING DATE: 2001-01-17
; Prior application data removed - consult PALM or file wrapper
; NUMBER OF SEQ ID NOS: 10231
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 6505
; LENGTH: 16163
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-764-891-6505

Query Match          39.6% Score 158.4; DB 3; Length 16163;
Best Local Similarity 73.9% Pred. No. 1.2e-37;
Matches 201; Conservative 0; Mismatches 71; Indels 0; Gaps 0;

QY 76 GCTGAATGCTGTATATCCAGCACTTCGGAGGCCAAGTGGCGGATCACCTGAGTCA 135
DB 8172 GGTTCACGCCCTGTATATCCAGCACTTCGGAGGCCAAGTGGCGGATCACCTGAGTCA 8231
QY 136 AGAGATGAGACCATCTCTGGCCAAATGTTAAACCCGCTCTTAACTAAATAACAAAA 195
DB 8232 GAGATTGAGATCACTCTGGCCAAATGTTAAACCCGCTCTTAACTAAATAACAAAA 8291
QY 196 ATAGCTGGGCAATGTGGACACACCTGTATGCTCCAGTCTACAGAGCCGAGAGATTGCGAG 255
DB 8292 CCAGCTGGGTGTGTGTGACACGCTTGTATTCAGGCTCTGAGCGGAGGTTGCGAG 8351
QY 256 TGAGCTGAGATTCGAGAGTGAAGCCGAATTCACAGATCAGAGTGAAGAGTGAACKC 315
DB 8352 TGAGCTGAGATTCGAGAGTGAAGCCGAATTCACAGATCAGAGTGAAGAGTGAACKC 8411
QY 316 CGTCTCAAAAACAACAACAAAAACAAAAA 347
DB 8412 GGAACAAAAACAAAAACAAAAACAAAAA 8443

RESULT 19
US-09-997-722-16
; Sequence 16, Application US/09997722
; Publication No. US20040072154A1
; GENERAL INFORMATION:
; APPLICANT: Engelhard, Eric
; TITLE OF INVENTION: NOVEL COMPOSITIONS AND METHODS FOR CANCER
; FILE REFERENCE: A-71171/RMS/DCF
; CURRENT APPLICATION NUMBER: US/09/997,722
; CURRENT FILING DATE: 2001-11-30
; Prior application data removed - See file wrapper or Palm
; Prior application number: US 09/747,377
; Prior application number: US 09/798,586
; Prior filing date: 2001-03-02
; NUMBER OF SEQ ID NOS: 301
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 16
; LENGTH: 96593
```

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QY 256 TGAGCTGATTCGAGAGTGAAGCCGAATTCACAGATCAGAGTGAAGAGTGAACKC 315
DB 8352 TGAGCTGATTCGAGAGTGAAGCCGAATTCACAGATCAGAGTGAAGAGTGAACKC 8411
QY 316 CGTCTCAAAAACAACAACAAAAACAAAAA 347
DB 8412 GGAACAAAAACAAAAACAAAAACAAAAA 8443

RESULT 18
US-10-091-414-270
; Sequence 270, Application US/10091414
; Publication No. US2003024461A1
; GENERAL INFORMATION:
; APPLICANT: Rosen et al.
; TITLE OF INVENTION: Nucleic Acids, Proteins, and Antibodies
; FILE REFERENCE: PA116C1
; CURRENT APPLICATION NUMBER: US/10/091,414
; CURRENT FILING DATE: 2002-03-07
; Prior application removed - See file wrapper or Palm
; NUMBER OF SEQ ID NOS: 392
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 270
; LENGTH: 16163
; TYPE: DNA
; ORGANISM: Homo sapiens
US-10-091-414-270

Query Match          39.6% Score 158.4; DB 7; Length 16163;
Best Local Similarity 73.9% Pred. No. 1.2e-37;
Matches 201; Conservative 0; Mismatches 71; Indels 0; Gaps 0;

QY 76 GCTGAATGCTGTATATCCAGCACTTCGGAGGCCAAGTGGCGGATCACCTGAGTCA 135
DB 8172 GGTTCACGCCCTGTATATCCAGCACTTCGGAGGCCAAGTGGCGGATCACCTGAGTCA 8231
QY 136 AGAGATGAGACCATCTCTGGCCAAATGTTAAACCCGCTCTTAACTAAATAACAAAA 195
DB 8232 GAGATTGAGATCACTCTGGCCAAATGTTAAACCCGCTCTTAACTAAATAACAAAA 8291
QY 196 ATAGCTGGGCAATGTGGACACACCTGTATGCTCCAGTCTACAGAGCCGAGAGATTGCGAG 255
DB 8292 CCAGCTGGGTGTGTGTGACACGCTTGTATTCAGGCTCTGAGCGGAGGTTGCGAG 8351
QY 256 TGAGCTGAGATTCGAGAGTGAAGCCGAATTCACAGATCAGAGTGAAGAGTGAACKC 315
DB 8352 TGAGCTGAGATTCGAGAGTGAAGCCGAATTCACAGATCAGAGTGAAGAGTGAACKC 8411
QY 316 CGTCTCAAAAACAACAACAAAAACAAAAA 347
DB 8412 GGAACAAAAACAAAAACAAAAACAAAAA 8443

RESULT 19
US-09-997-722-16
; Sequence 16, Application US/09997722
; Publication No. US20040072154A1
; GENERAL INFORMATION:
; APPLICANT: Engelhard, Eric
; TITLE OF INVENTION: NOVEL COMPOSITIONS AND METHODS FOR CANCER
; FILE REFERENCE: A-71171/RMS/DCF
; CURRENT APPLICATION NUMBER: US/09/997,722
; CURRENT FILING DATE: 2001-11-30
; Prior application data removed - See file wrapper or Palm
; Prior application number: US 09/747,377
; Prior application number: US 09/798,586
; Prior filing date: 2001-03-02
; NUMBER OF SEQ ID NOS: 301
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 16
; LENGTH: 96593
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; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-997-722-16

Query Match      39.6%; Score 158.4; DB 3; Length 96593;
Best Local Similarity 75.0%; Pred. No. 2.5e-37;
Matches 216; Conservative 1; Mismatches 57; Indels 14; Gaps 1;

OY 81 ATGCTGTAAATCCAGACCTTCGGAGGCGCAAGTGGCGGATCACTGAGTCAAGAGA 140
DB 18217 ACGCTGTAAATCCAGACCTTCGGAGGCGCAAGTGGCGGATTCCTGAGTCAAGAGA 18276
OY 141 TCGAGACCATCTGCGCCCAACATGTTGAATACCCCGCTCTTACTTAAATAATCAAAAAATATAGC 200
DB 18277 TCAAGATACAGCTGCGCCCAACATGTTGAATACCCCGCTCTTACTTAAATAATCAAAAAATATAGC 18336
OY 201 TGGGCAATGTGGCAACACCTGTAGTCCAGCTACTCAGAGCGGAGATTGCAATGAGC 260
DB 18337 TGGGCATGTGGCGGCACTGTAGTCCAGCTACTCAGAGCGGAGATTGCAATGAGC 18396
OY 261 TGAAGTCCAG-----AGTACCCCAATCAAGATCAGAGTGAAGCAGA 306
DB 18397 CTGTGAACCCAGCGCGGAGGTTCAGATGAGCCGAGATCGACCACTTGGGTGACAGAGC 18456
OY 307 GTGAGACKCCGCTCTCAAAAACACAAACAAAAAACCAATTAAG 354
DB 18457 GTGAGACTTCACTCTCAAAAACAAAAAACCAATTAAG 18504

RESULT 20
US-10-301-480-603905/c
; Sequence 603905, Application US/10301480
; Publication No. US20060057564A1
; GENERAL INFORMATION:
; APPLICANT: Wang, David G.
; TITLE OF INVENTION: Identification and Mapping of Single Nucleotide Polymorphisms
; FILE REFERENCE: 108827.137
; CURRENT FILING DATE: 2002-11-21
; PRIOR APPLICATION NUMBER: US 10/215,598
; PRIOR FILING DATE: 2002-08-09
; PRIOR APPLICATION NUMBER: US 60/311,695
; PRIOR FILING DATE: 2001-08-10
; NUMBER OF SEQ ID NOS: 1226818
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 603905
; LENGTH: 997
; TYPE: DNA
; ORGANISM: Homo sapien
US-10-301-480-603905

Query Match      39.6%; Score 158.2; DB 12; Length 997;
Best Local Similarity 78.5%; Pred. No. 4.7e-38;
Matches 205; Conservative 0; Mismatches 48; Indels 8; Gaps 1;

OY 81 ATGCTGTAAATCCAGACCTTCGGAGGCGCAAGTGGCGGATTCAGTCAAGAGA 140
DB 733 ATGCTGTAAATCCAGACCTTCGGAGGCGCAAGTGGCGGATTCAGTCAAGAGA 674
OY 141 TCGAGACCATCTGCGCCCAACATGTTGAATACCCCGCTCTTACTTAAATAATCAAAAAATATAGC 200
DB 673 TCAAGATACAGCTGCGCCCAACATGTTGAATACCCCGCTCTTACTTAAATAATCAAAAAATATAGC 614
OY 201 TGGGCAATGTGGCAACACCTGTAGTCCAGCTACTCAGAGCGGAGATTGCAATGAGC 260
DB 613 TGGGCATGTGGCGGCACTGTAGTCCAGCTACTCAGAGCGGAGATTGCAATGAGC 554
OY 261 TGAAGTCCAGAGTGAAGCGCAATCAAGATCAGAGTGAAGTGAAGCAGCAGTCT 320
DB 553 TGAAGTTG-----TGCACCTGCACTCCAGCTGGGTGACAAAGACAAACACTGTCT 502
OY 321 CAAAAACAACAACAAAAACA 341
DB 501 CAAAAACAAAAACAAAAAGACA 481
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DB 501 CAAAAACAAAAACAAAAAGACA 481

RESULT 21
US-10-301-480-1217314/c
; Sequence 1217314, Application US/10301480
; Publication No. US20060057564A1
; GENERAL INFORMATION:
; APPLICANT: Wang, David G.
; TITLE OF INVENTION: Identification and Mapping of Single Nucleotide Polymorphisms
; FILE REFERENCE: 108827.137
; CURRENT FILING DATE: 2002-11-21
; PRIOR APPLICATION NUMBER: US/10/301,480
; PRIOR FILING DATE: 2002-11-21
; PRIOR APPLICATION NUMBER: US 10/215,598
; PRIOR FILING DATE: 2002-08-09
; PRIOR APPLICATION NUMBER: US 60/311,695
; PRIOR FILING DATE: 2001-08-10
; NUMBER OF SEQ ID NOS: 1226818
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 1217314
; LENGTH: 997
; TYPE: DNA
; ORGANISM: Homo sapien
US-10-301-480-1217314

Query Match      39.6%; Score 158.2; DB 12; Length 997;
Best Local Similarity 78.5%; Pred. No. 4.7e-38;
Matches 205; Conservative 0; Mismatches 48; Indels 8; Gaps 1;

OY 81 ATGCTGTAAATCCAGACCTTCGGAGGCGCAAGTGGCGGATTCAGTCAAGAGA 140
DB 733 ATGCTGTAAATCCAGACCTTCGGAGGCGCAAGTGGCGGATTCAGTCAAGAGA 674
OY 141 TCGAGACCATCTGCGCCCAACATGTTGAATACCCCGCTCTTACTTAAATAATCAAAAAATATAGC 200
DB 673 TCAAGATACAGCTGCGCCCAACATGTTGAATACCCCGCTCTTACTTAAATAATCAAAAAATATAGC 614
OY 201 TGGGCAATGTGGCAACACCTGTAGTCCAGCTACTCAGAGCGGAGATTGCAATGAGC 260
DB 613 TGGGCATGTGGCGGCACTGTAGTCCAGCTACTCAGAGCGGAGATTGCAATGAGC 554
OY 261 TGAAGTCCAGAGTGAAGCGCAATCAAGATCAGAGTGAAGTGAAGCAGCAGTCT 320
DB 553 TGAAGTTG-----TGCACCTGCACTCCAGCTGGGTGACAAAGACAAACACTGTCT 502
OY 321 CAAAAACAACAACAAAAACA 341
DB 501 CAAAAACAAAAACAAAAAGACA 481

RESULT 22
US-10-301-480-577516
; Sequence 577516, Application US/10301480
; Publication No. US20060057564A1
; GENERAL INFORMATION:
; APPLICANT: Wang, David G.
; TITLE OF INVENTION: Identification and Mapping of Single Nucleotide Polymorphisms
; FILE REFERENCE: 108827.137
; CURRENT FILING DATE: 2002-11-21
; PRIOR APPLICATION NUMBER: US/10/301,480
; PRIOR FILING DATE: 2002-11-21
; PRIOR APPLICATION NUMBER: US 10/215,598
; PRIOR FILING DATE: 2002-08-09
; PRIOR APPLICATION NUMBER: US 60/311,695
; PRIOR FILING DATE: 2001-08-10
; NUMBER OF SEQ ID NOS: 1226818
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 577516
; LENGTH: 635
; TYPE: DNA
; ORGANISM: Homo sapien
```

US-10-301-480-577516

Query Match 39.4%; Score 157.4; DB 12; Length 635;

Best Local Similarity 77.2%; Pred. No. 6.9e-38;

Matches 206; Conservative 1; Mismatches 52; Indels 8; Gaps 1;

QY 81 ATGCCGTATATCCAGCACTTCGGAGGCCAAGTGGGGGATACCTGAGTCAAGAGA 140
 DB 252 AGCGCTGTATATCCAGCACTTCGGAGGCCAAGTGGGGGATACCTGAGTCAAGAGA 311
 QY 141 TCGAGCCATCTCTGGCCAACTGTGTAAACCCCGTCTTACTTAAATAAATAAATAGC 200
 DB 312 TCAAGACCAAGCTGGCCAAATGTGTAAACCCCATCTCTAATAAATAAATAAATAGC 371
 QY 201 TGGGATGGTGGAACACCTGTAGTCCAGCTACTCAGAGCCGGAATTGCAATGAGC 260
 DB 372 TGGGATGTATGGGGGTACTGTATCCAGCTACTCAGAGCCGGAATTGCGGTGAGC 431
 QY 261 TGAGATCGCAGATGAGCCGAATCAAGATCAAGATGAGCAGTGAACKCCGCTCT 320
 DB 432 CGAGATCG-----TGCATTGCAACCCAGCTTGGGCAAGAGCGAAACTCTGTCT 483
 QY 321 CAAAAACAACAACAAAAA 347
 DB 484 CAAAAAAGAAAAAAGAAAA 510

RESULT 23

US-10-301-480-1190925

Sequence 1190925, Application US/10301480

Publication No. US20060057564A1

GENERAL INFORMATION:

APPLICANT: Wang, David G.

TITLE OF INVENTION: Identification and Mapping of Single Nucleotide Polymorphisms

FILE REFERENCE: 108827.137

CURRENT APPLICATION NUMBER: US/10/301,480

CURRENT FILING DATE: 2002-11-21

PRIOR APPLICATION NUMBER: US 10/215,598

PRIOR FILING DATE: 2002-08-09

PRIOR APPLICATION NUMBER: US 60/311,695

PRIOR FILING DATE: 2001-08-10

NUMBER OF SEQ ID NOS: 1226818

SOFTWARE: FastSeq for Windows Version 4.0

SEQ ID NO 1190925

LENGTH: 635

TYPE: DNA

ORGANISM: Homo sapien

US-10-301-480-1190925

Query Match 39.4%; Score 157.4; DB 12; Length 635;

Best Local Similarity 77.2%; Pred. No. 6.9e-38;

Matches 206; Conservative 1; Mismatches 52; Indels 8; Gaps 1;

QY 81 ATGCCGTATATCCAGCACTTCGGAGGCCAAGTGGGGGATACCTGAGTCAAGAGA 140
 DB 252 AGCGCTGTATATCCAGCACTTCGGAGGCCAAGTGGGGGATACCTGAGTCAAGAGA 311
 QY 141 TCGAGCCATCTCTGGCCAACTGTGTAAACCCCGTCTTACTTAAATAAATAAATAGC 200
 DB 312 TCAAGACCAAGCTGGCCAAATGTGTAAACCCCATCTCTAATAAATAAATAAATAGC 371
 QY 201 TGGGATGGTGGAACACCTGTAGTCCAGCTACTCAGAGCCGGAATTGCAATGAGC 260
 DB 372 TGGGATGTATGGGGGTACTGTATCCAGCTACTCAGAGCCGGAATTGCGGTGAGC 431
 QY 261 TGAGATCGCAGATGAGCCGAATCAAGATCAAGATGAGCAGTGAACKCCGCTCT 320
 DB 432 CGAGATCG-----TGCATTGCAACCCAGCTTGGGCAAGAGCGAAACTCTGTCT 483
 QY 321 CAAAAACAACAACAAAAA 347
 DB 484 CAAAAAAGAAAAAAGAAAA 510

RESULT 24

US-10-027-632-98771/c

Sequence 98771, Application US/10027632

Publication No. US20020198371A1

GENERAL INFORMATION:

APPLICANT: Wang, David G.

TITLE OF INVENTION: Identification and Mapping of Single Nucleotide

FILE REFERENCE: 108827.129

CURRENT APPLICATION NUMBER: US/10/027,632

CURRENT FILING DATE: 2002-04-30

PRIOR APPLICATION NUMBER: US 60/218,006

PRIOR FILING DATE: 2000-07-12

PRIOR APPLICATION NUMBER: US 60/198,676

PRIOR FILING DATE: 2000-04-20

PRIOR APPLICATION NUMBER: US 60/193,483

PRIOR FILING DATE: 2000-03-29

PRIOR APPLICATION NUMBER: US 60/185,218

PRIOR FILING DATE: 2000-02-24

PRIOR APPLICATION NUMBER: US 60/167,363

PRIOR FILING DATE: 1999-11-23

PRIOR APPLICATION NUMBER: US 60/156,358

PRIOR FILING DATE: 1999-09-28

PRIOR APPLICATION NUMBER: US 60/146,002

PRIOR FILING DATE: 1999-08-09

NUMBER OF SEQ ID NOS: 325720

SOFTWARE: FastSeq for Windows Version 4.0

SEQ ID NO 98771

LENGTH: 1963

TYPE: DNA

ORGANISM: Human

US-10-027-632-98771

Query Match 38.9%; Score 155.4; DB 6; Length 1963;

Best Local Similarity 72.6%; Pred. No. 4.5e-37;

Matches 217; Conservative 1; Mismatches 72; Indels 9; Gaps 1;

QY 81 ATGCCGTATATCCAGCACTTCGGAGGCCAAGTGGGGGATACCTGAGTCAAGAGA 140
 DB 463 AGCGCTGTATATCCAGCACTTCGGAGGCCAAGTGGGGGATACCTGAGTCAAGAGA 404
 QY 141 TCGAGCCATCTCTGGCCAACTGTGTAAACCCCGTCTTACTTAAATAAATAAATAGC 200
 DB 403 TCGAGCCATCTCTGGCCAACTGTGTAAACCCCGTCTTACTTAAATAAATAAATAGC 344
 QY 201 TGGGATGGTGGAACACCTGTAGTCCAGCTACTCAGAGCCGGAATTGCAATGAGC 260
 DB 343 CGGATGTGTGGGGGTACTGTATCCAGCTACTCAGAGCCGGAATTGCAATGAGC 284
 QY 261 TGAGATCGCAGATGAGCCGAATCAAGATCAAGATGAGCAGTGAACKCCGCTCT 320
 DB 283 CGAGATCGCTTGTGACTTCCAGCTTGGGGA-----CAGACGAGACTCTGTCT 233
 QY 321 CAAAAACAACAACAAAAA 347
 DB 232 CAAACAATAACAACAAAAA 379

RESULT 25

US-10-027-632-98771/c

Sequence 98771, Application US/10027632

Publication No. US20030204075A9

GENERAL INFORMATION:

APPLICANT: Wang, David G.

TITLE OF INVENTION: Identification and Mapping of Single Nucleotide

FILE REFERENCE: 108827.129

CURRENT APPLICATION NUMBER: US/10/027,632

CURRENT FILING DATE: 2002-04-30

PRIOR APPLICATION NUMBER: US 60/218,006

PRIOR FILING DATE: 2000-07-12

Query Match	38.9%	Score 155.4	DB 6	Length 31898
Best Local Similarity	72.6%	Pred. No. 1.4e-36		
Matches 217, Conservative		1, Mismatches 9		Gaps 1

Qy 288 AGATCAGAGTGAAGCAGAGTGAGACKCCGTCCTCAAAAAACAACAACAAAAACAAAAA 347
| | | | | | | | | | | | | | | | | | | | | | : | | | |
Db 374 TCCACCTCGGAGACA-CAGAGGAGACTGTGTCTCAAAAAAAAAAAAAAAAAAAAAA 316


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RESULT 29
US-10-301-480-609460/c
: Sequence 609460, Application US/10301480
: Publication No. US20060057564A1
: GENERAL INFORMATION:
: APPLICANT: Wang, David G.
: TITLE OF INVENTION: Identification and Mapping of Single Nucleotide Polymorphisms
: TITLE OF INVENTION: in the Human Genome
: FILE REFERENCE: 108827.137

```

Query Match	38.4%	Score 153.6	DB 12	Length 977
Best Local Similarity	70.3%	Pred. No. 1.2e-36		
Matches 218	Conservative 1	Mismatches 90	Indels 1	Gaps 1

OY	48	AGACATTGTCAAGCCGATGACACTGGCTGAATGCTTAAATCCACAGCTTCGGAG	107
436	AGAAACAGTGTGACGCGCGGTGCGGTGCGGCACACTCTAATCCACAGACTTTGGAG	377	

QY 108 GCCAAGTGGGGGATCATCTGAGTCAAGAGATCGAGACCATCTCGGCCAATGTGGA 167
DB 376 GCCAAGTGGGGGATCGCTCGAGATCGAGATCGAGACCATCTCGGCCAATGTGGA 317
QY 168 AACCCGCTTCTTAATAAATAAATAAATAAATAAATAAATAAATAAATAAATAAATAA 227
DB 316 AACCCGCTTCTTAATAAATAAATAAATAAATAAATAAATAAATAAATAAATAAATAA 257
QY 228 CCAGCTACTCAGAGCGGAGATTCAGTGAAGTGCAGAGTGCAGAGTGCAGAGTGCAG 287
DB 256 CCAGCTACTCAGAGCGGAGATTCAGTGAAGTGCAGAGTGCAGAGTGCAGAGTGCAG 197
QY 288 AGATCAGAGTGCAGAGTGCAGAGTGCAGAGTGCAGAGTGCAGAGTGCAGAGTGCAG 347
DB 196 TCCAGCTCGGAGAG-CAGAGGAGAGCTCTGTCTCAAAAAAAAAAAAAAAAAAAAAA 138
QY 348 CCATAGACA 357
DB 137 GAAGAAAAAA 128

RESULT 31
US-10-995-561-13274/c
; Sequence 13274, Application US/10995561
; Publication No. US20050272054A1
; GENERAL INFORMATION:
; APPLICANT: CARILL, Michele et al.
; TITLE OF INVENTION: GENETIC POLYMORPHISMS ASSOCIATED WITH
; TITLE OF INVENTION: CARDIOVASCULAR DISORDERS AND DRUG RESPONSE, METHODS OF
; FILE REFERENCE: CLO01559
; CURRENT APPLICATION NUMBER: US/10/995,561
; CURRENT FILING DATE: 2004-11-24
; NUMBER OF SEQ ID NOS: 85702
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 13274
; LENGTH: 415117
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: misc feature
; LOCATION: (1) ..(415117)
; OTHER INFORMATION: n = A,T,C or G, or insertion/deletion polymorphism (see Tables 1-
US-10-995-561-13274

Query Match 38.4%; Score 153.4; DB 10; Length 415117;
Best Local Similarity 68.5%; Pred. No. 1.6e-35;
Matches 224; Conservative 2; Mismatches 98; Indels 3; Gaps 1;
QY 39 ATATTAAATGACATTTGTCAGAGCCGAGCATGCTGCTGAATTCCTGTAATCCCGACA 98
DB 228512 ATATTAAATGACATTTGTCAGAGCCGAGCATGCTGCTGAATTCCTGTAATCCCGACA 228453
QY 99 CTTGGGAGGCGCAAGGTGGCGGATCACCTGAGTCAAGAGTGCAGAGTGCAGAGTGCAG 158
DB 228452 CTTGGAGGCGCGAGGTGGGATGATCTTGAAGTCAAGAGTGCAGAGTGCAGAGTGCAG 228393
QY 159 ACATGTGTAACCCCGTCTTTACTTAATAAATAAATAAATAAATAAATAAATAAATAA 218
DB 228392 ACATGTGTAACCCCGTCTTTACTTAATAAATAAATAAATAAATAAATAAATAAATAA 228333
QY 219 CCGTGAAGTCCAGCTACTCAGAGCGGAGATTCAGAGTGCAGAGTGCAGAGTGCAG 278
DB 228332 CCGTGAAGTCCAGCTACTCAGAGCGGAGATTCAGAGTGCAGAGTGCAGAGTGCAG 228273
QY 279 CGAAATCAGATGATGAGAGTGCAGAGTGCAGAGTGCAGAGTGCAGAGTGCAGAGTGCAG 338
DB 228272 CTCCAGCTCGGCGGAGAGG---GGAGACTCGGTCTCAACCAAAAAAAAAAAAAAGTAA 228216
QY 339 ACAAAAAAACCATTAAGACATTTGTCAT 365
DB 228215 AGAAAGTAAATTAATTAATTCAT 228189

RESULT 32
US-10-301-480-598906
; Sequence 598906, Application US/10301480
; Publication No. US20060057564A1
; GENERAL INFORMATION:
; APPLICANT: Wang, David G.
; TITLE OF INVENTION: Identification and Mapping of Single Nucleotide Polymorphisms
; FILE REFERENCE: 108827.137
; CURRENT APPLICATION NUMBER: US/10/301,480
; CURRENT FILING DATE: 2002-11-21
; PRIOR APPLICATION NUMBER: US 10/215,598
; PRIOR FILING DATE: 2002-08-09
; PRIOR APPLICATION NUMBER: US 60/311,695
; PRIOR FILING DATE: 2001-08-10
; NUMBER OF SEQ ID NOS: 1226818
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 598906
; LENGTH: 984
; TYPE: DNA
; ORGANISM: Homo sapien
US-10-301-480-598906

Query Match 38.3%; Score 153; DB 12; Length 984;
Best Local Similarity 71.5%; Pred. No. 1.9e-36;
Matches 201; Conservative 0; Mismatches 80; Indels 0; Gaps 0;
QY 81 ATGCTGTAATCCAGCAGCTTCGGAGGCGCAAGTGGCGGATCACCTGAGTCAAGAGA 140
DB 225 ATGCTGTAATCCAGCAGCTTCGGAGGCGCAAGTGGCGGATCACCTGAGTCAAGAGA 284
QY 141 TCGAGACCATCTCTGCGCAATGAGTGAACCCGCTCTTTACTTAATAAATAAATAAATAA 200
DB 285 TCGAGACCATCTCTGCGCAATGAGTGAACCCGCTCTTTACTTAATAAATAAATAAATAA 344
QY 201 TGGGATGATGGGACACACACCTGTAGTCCAGCTACTCAGAGGCGGAGATTGAGTGAAGC 260
DB 345 CAGGATGATGGGACACACCTGTAGTCCAGCTACTCAGAGGCGGAGATTGAGTGAAGTGA 404
QY 261 TGAGATCGAGAGTGAAGCCGAAATCAGAGTCAAGAGTGAAGAGTGAAGTGAAGTGAAG 320
DB 405 CGTAGGTGAGAGTGAAGTGAAGTGAAGTGAAGTGAAGTGAAGTGAAGTGAAGTGAAGTGA 464
QY 321 CAAAAACAACAACAAAAAATAAATAAATAAATAAATAAATAAATAAATAAATAAATAA 361
DB 465 CTCTGTCTCAAAAAAAAAAAAAAAAAATAAATAAATAAATAAATAAATAAATAAATAA 505

RESULT 33
US-10-301-480-1212315
; Sequence 1212315, Application US/10301480
; Publication No. US20060057564A1
; GENERAL INFORMATION:
; APPLICANT: Wang, David G.
; TITLE OF INVENTION: Identification and Mapping of Single Nucleotide Polymorphisms
; FILE REFERENCE: 108827.137
; CURRENT APPLICATION NUMBER: US/10/301,480
; CURRENT FILING DATE: 2002-11-21
; PRIOR APPLICATION NUMBER: US 10/215,598
; PRIOR FILING DATE: 2002-08-09
; PRIOR APPLICATION NUMBER: US 60/311,695
; PRIOR FILING DATE: 2001-08-10
; NUMBER OF SEQ ID NOS: 1226818
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 1212315
; LENGTH: 984
; TYPE: DNA
; ORGANISM: Homo sapien
US-10-301-480-1212315

```

Query Match          38.3%; Score 153; DB 12; Length 984;
Best Local Similarity 71.5%; Pred. No. 1.9e-36;
Matches 201; Conservative 0; Mismatches 80; Indels 0; Gaps 0;

QY 81 ATGCTGTATATCCAGCACTTGGGAGGCCAAGTGGGGATCACCTGAGTCAAGAGA 140
DB 225 ATGCTGTATATCCAGCACTTGGGAGGCCAAGTGGGGATCACCTGAGTCAAGAGA 284
QY 141 TCGAGACCATCTGGCCAAATGTAATCCCGCTTTACTTAAATAATCAAAAAATAGC 200
DB 225 TCGAGACCATCTGGCCAAATGTAATCCCGCTTTACTTAAATAATCAAAAAATAGC 344
QY 201 TGGGCAATGTGGCAACAACCTGTAGTCCAGCTACTCAGAGCCGAGATTGCAAGTAC 260
DB 345 CAGGATGATGTGGGCAACTGTATATCCAGCTACTCAGAGAGGCTGAGACAGAGATTA 404
QY 261 TGAGATCCAGAGTGAACCGGAATCAAGATCAAGAGTGAAGCAAGTGAAGCCKCGTCT 320
DB 405 CGTGAAGTTCAGTGAAGTGAATCAAGCACTGCACTCAGCTTGAAGGACAGAGATG 464
QY 321 CAAAAACAACAACAAAAAACAATAAGCATTTGT 361
DB 465 CTCTGTCTCAAAAAAATAAATTAAGTGGGCAATAGT 505

RESULT 34
US-09-764-891-7647/c
; Sequence 7647, Application US/09764891
; Publication No. US20030077808A1
; GENERAL INFORMATION:
; APPLICANT: Rosen et al.
; TITLE OF INVENTION: Nucleic Acids, Proteins, and Antibodies
; FILE REFERENCE: PC006
; CURRENT APPLICATION NUMBER: US/09/764,891
; CURRENT FILING DATE: 2001-01-17
; Prior application data removed - consult PALM or file wrapper
; NUMBER OF SEQ ID NOS: 10231
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 7647
; LENGTH: 4388
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-764-891-7647

Query Match          38.1%; Score 152.4; DB 3; Length 4388;
Best Local Similarity 76.5%; Pred. No. 5.3e-36;
Matches 202; Conservative 1; Mismatches 52; Indels 9; Gaps 1;

QY 81 ATGCTGTATATCCAGCACTTGGGAGGCCAAGTGGGGATCACCTGAGTCAAGAGA 140
DB 1275 AAGCTGTATATCCAGCACTTGGGAGGCCAAGTGGGGATCACCTGAGTCAAGAGA 1216
QY 141 TCGAGACCATCTGGCCAAATGTAATCCCGCTTTACTTAAATAATCAAAAAATAGC 200
DB 1215 TCGAGACCATCTGGCCAAATGTAATCCCGCTTTACTTAAATAATCAAAAAATAGC 1156
QY 201 TGGGCAATGTGGCAACAACCTGTAGTCCAGCTACTCAGAGCCGAGATTGCAAGTAC 260
DB 1155 CGCAGTGTGGGCGCGCGCTGTATCCAGCTACTCAGAGGCGGAGTGGCAGCGAGC 1096
QY 261 TGAGATCCAGAGTGAACCGGAATCAAGATCAAGAGTGAAGCAAGTGAAGCCKCGTCT 320
DB 1095 AGAGATGTGCTGCAATTCAGCTCTGAGGCGA-----CACAGGAGATCTCGCT 1045
QY 321 CAAAAACAACAACAAAAAACAATAAGCATTTGT 344
DB 1044 CAAAAAATAAATAAATAAATAAATAAATAA 1021

RESULT 35
US-09-967-768A-314/c
; Sequence 314, Application US/09967768A
; Patent No. US2002015087A1

```

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; GENERAL INFORMATION:
; APPLICANT: Augustus, Meena
; TITLE OF INVENTION: Cancer Gene Determination and Therapeutic Screening Using Signat
; TITLE OF INVENTION: Sets
; FILE REFERENCE: 689290-72
; CURRENT APPLICATION NUMBER: US/09/967,768A
; CURRENT FILING DATE: 2001-09-28
; PRIOR APPLICATION NUMBER: US/60/236,109
; PRIOR FILING DATE: 2000-09-28
; PRIOR APPLICATION NUMBER: US/60/236,034
; PRIOR FILING DATE: 2000-09-28
; PRIOR APPLICATION NUMBER: US/60/236,111
; PRIOR FILING DATE: 2000-09-28
; NUMBER OF SEQ ID NOS: 325
; SOFTWARE: PatentIn version 3.0
; SEQ ID NO 314
; LENGTH: 174424
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-967-768A-314

Query Match          38.0%; Score 152; DB 3; Length 174424;
Best Local Similarity 71.8%; Pred. No. 3e-35;
Matches 214; Conservative 0; Mismatches 80; Indels 4; Gaps 1;

QY 6 TACTAGCCATGTGCTGGGCCATGGGAACCAATATTAAATPAGACATTTGCGCCAGG 65
DB 48438 TCTTCAGGTGTCTGTGTATCTTGTGTCTTCTCATATTTAAAGATGGGCGAGCTGGG 48379
QY 66 CATGACACTGGCTGAATGCTGTATATCCAGCACTTGGGAGGCCAAGTGGGCGGATCA 125
DB 48378 CATGCA-----GCTCATGCTGTATATCCAGCACTTGGGAGGCCAAGTGGGCGGATCA 48323
QY 126 CTTGAGGTCAAGAGATCGAGACATCTTGGCCAAATGTAATCCCGCTTTACTTAA 185
DB 48322 CTTGAGGTCAAGAGATCGAGACATCTTGGCCAAATGTAATCCCGCTTTACTTAA 48263
QY 186 AATACAAAAAATAGTGGGCGATGTGGCAACACCTGTAGTCCAGCTACTCAGAGCGG 245
DB 48262 AATACAAAAAATAGTGGGCGATGTGGCAACACCTGTAGTCCAGCTACTCAGAGCGG 48203
QY 246 GAGATGAGTGAAGTGAATCGAGAGTGAAGCGGAATATCAAGATCAAGAGTGAAGC 303
DB 48202 GAGGAGAGATGCTTGAATCCGAGAGTGAAGTGAAGTGAAGTGAAGTGAAGTGAAG 48145

RESULT 36
US-09-960-706-969/c
; Sequence 969, Application US/09960706
; Publication No. US20030134280A1
; GENERAL INFORMATION:
; APPLICANT: Munger, William B.
; TITLE OF INVENTION: Identifying Drugs for and Diagnosis of Benign Prostatic Hyperplas
; FILE REFERENCE: 44921-5029-01US
; CURRENT APPLICATION NUMBER: US/09/960,706
; CURRENT FILING DATE: 2001-09-24
; PRIOR APPLICATION NUMBER: 60/223,323
; PRIOR FILING DATE: 2000-08-07
; PRIOR APPLICATION NUMBER: 09/873,319
; PRIOR FILING DATE: 2001-06-05
; NUMBER OF SEQ ID NOS: 1124
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 969
; LENGTH: 174424
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; OTHER INFORMATION: Genbank Accession No. US20030134280A1 US2112
US-09-960-706-969

Query Match          38.0%; Score 152; DB 3; Length 174424;
Best Local Similarity 71.8%; Pred. No. 3e-35;

```

```
Matches 214; Conservative 0; Mismatches 80; Indels 4; Gaps 1;
QY 6 TACTCAGCCATGTCCTGGCCATGGAAACCAATATTATTAAGACATTGTCAAGCCAG 65
Db 48438 TCCCTCAGGTGTCTGGTATCTTGTGTCTTCTCATATTAAAGAGGGCAGGCTGGG 48379
QY 66 CATGACACTGGCTGAATGCTGTATCCCAAGCACTTGGGGGCGAAGTGGGGGATCA 125
Db 48378 CATGGCA---GCTCATGCTCTGTATCTCCACACTTTGGAGGCGCAAGTGGGGATCA 48323
QY 126 CTTGAGGTCAAGAGATCGAGACCATCTGGCCAAATGTTGMAAACCCTGTTACTAAA 185
Db 48322 CTTGAGGTCAAGAGATCAAGTCCATCTCGCCCAATGTTGMAAACCCTGTTACTAAA 48263
QY 186 AATACAAAAATTAAGCTGGGCGATGGTGACACACCTGTAGTCCAGCTACTCAGAGCCG 245
Db 48262 AATACAAAAATTAAGCTGGGCGATGGTGACACACCTGTAGTCCAGCTACTTGGAGGCT 48203
QY 246 GAGATTGCAGTGAAGCTGATCGAGAGTGAGCCGAATATCAGATCAAGATGAGC 303
Db 48202 GAGGCAAGAGATCGCTTGAATCCGGAGGTGAGATTGCACTGAGCCGAGATTGTGC 48145

RESULT 37
US-10-843-641A-6459/c
; Sequence 6459, Application US/10843641A
; Publication No. US20050064454A1
; GENERAL INFORMATION:
; APPLICANT: Avalon Pharmaceuticals, Inc.
; TITLE OF INVENTION: Cancer Gene Determination and Therapeutic Screening Using
; FILE REFERENCE: 689290-189
; CURRENT APPLICATION NUMBER: US/10/843, 641A
; PRIOR FILING DATE: 2004-05-12
; PRIOR APPLICATION NUMBER: US/09/873,367
; PRIOR FILING DATE: 2001-06-05
; PRIOR APPLICATION NUMBER: US/09/954,531
; PRIOR FILING DATE: 2001-09-18
; PRIOR APPLICATION NUMBER: US/09/954,456
; PRIOR FILING DATE: 2001-09-25
; PRIOR APPLICATION NUMBER: US/09/962,436
; PRIOR FILING DATE: 2001-09-25
; PRIOR APPLICATION NUMBER: US/09/962,832
; PRIOR FILING DATE: 2001-09-25
; PRIOR APPLICATION NUMBER: US/09/964,824
; PRIOR FILING DATE: 2001-09-27
; PRIOR APPLICATION NUMBER: US/09/967,768
; PRIOR FILING DATE: 2001-09-28
; PRIOR APPLICATION NUMBER: US/09/968,007
; PRIOR FILING DATE: 2001-10-02
; PRIOR APPLICATION NUMBER: US/09/969,347
; PRIOR FILING DATE: 2001-10-02
; PRIOR APPLICATION NUMBER: US/09/969,708
; PRIOR FILING DATE: 2001-10-03
; Remaining Prior Application data removed - See File Wrapper or PALM.
; NUMBER OF SEQ ID NOS: 8447
; SOFTWARE: PatentIn version 3.0
; SEQ ID NO 6459
; LENGTH: 174424
; TYPE: DNA
; ORGANISM: Homo sapiens
US-10-843-641A-6459

Query Match 38.0%; Score 152; DB 10; Length 174424;
Best Local Similarity 71.8%; Pred. No. 3e-35;
Matches 214; Conservative 0; Mismatches 80; Indels 4; Gaps 1;
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```
Db 48378 CATGGCA---GCTCATGCTCTGTATATCCAAACCTTTGGAGGCGAAGTGGGCGATCA 48323
QY 126 CTTGAGGTCAAGAGATGAGAACCATCTGGCCAAATGTTGMAAACCCTGTTACTAAA 185
Db 48322 CTTGAGGTCAAGAGATGAGAACCATCTGGCCAAATGTTGMAAACCCTGTTACTAAA 48263
QY 186 AATACAAAAATTAAGCTGGGCGATGGTGACACACCTGTAGTCCAGCTACTCAGAGCCG 245
Db 48262 AATACAAAAATTAAGCTGGGCGATGGTGACACACCTGTAGTCCAGCTACTTGGAGGCT 48203
QY 246 GAGATTGCAGTGAAGCTGATCGAGAGTGAGCCGAATATCAGATCAAGATGAGC 303
Db 48202 GAGGCAAGAGATCGCTTGAATCCGGAGGTGAGATTGCACTGAGCCGAGATTGTGC 48145

RESULT 38
US-10-723-860-2392/c
; Sequence 2392, Application US/10723860
; Publication No. US20040253606A1
; GENERAL INFORMATION:
; APPLICANT: Aziz, Natasha
; APPLICANT: Ginsburg, Wendy M.
; APPLICANT: Zlotnik, Albert
; TITLE OF INVENTION: Methods of Diagnosis of Soft Tissue Sarcoma, Compositions &
; TITLE OF INVENTION: Methods for Screening for Soft Tissue Sarcoma Modulators
; FILE REFERENCE: 05882,0193,NPUS01
; CURRENT APPLICATION NUMBER: US/10/723, 860
; CURRENT FILING DATE: 2003-11-26
; PRIOR APPLICATION NUMBER: 60/429,739
; PRIOR FILING DATE: 2002-11-26
; NUMBER OF SEQ ID NOS: 8393
; SOFTWARE: PatentIn version 3.2
; SEQ ID NO 2392
; LENGTH: 181343
; TYPE: DNA
; ORGANISM: Homo sapiens
US-10-723-860-2392

Query Match 38.0%; Score 152; DB 9; Length 181343;
Best Local Similarity 71.8%; Pred. No. 3.1e-35;
Matches 214; Conservative 0; Mismatches 80; Indels 4; Gaps 1;
QY 6 TACTCAGCCATGTCCTGGCCATGGAAACCAATATTATTAAGACATTGTCAAGCCAG 65
Db 48439 TCCCTCAGGTGTCTGGTATCTTGTGTCTTCTCATATTAAAGAGGGCAGGCTGGG 48380
QY 66 CATGACACTGGCTGAATGCTGTATATCCCAAGCACTTGGGAGGCGCAAGTGGGGGATCA 125
Db 48379 CATGGCA---GCTCATGCTCTGTATATCCCAACCTTTGGAGGCGCAAGTGGGGGATCA 48324
QY 126 CTTGAGGTCAAGAGATGAGAACCATCTGGCCAAATGTTGMAAACCCTGTTACTAAA 185
Db 48322 CTTGAGGTCAAGAGATGAGAACCATCTGGCCAAATGTTGMAAACCCTGTTACTAAA 48264
QY 186 AATACAAAAATTAAGCTGGGCGATGGTGACACACCTGTAGTCCAGCTACTCAGAGCCG 245
Db 48263 AATACAAAAATTAAGCTGGGCGATGGTGACACACCTGTAGTCCAGCTACTTGGAGGCT 48204
QY 246 GAGATTGCAGTGAAGCTGATCGAGAGTGAGCCGAATATCAGATCAAGATGAGC 303
Db 48203 GAGGCAAGAGATCGCTTGAATCCGGAGGTGAGATTGCACTGAGCCGAGATTGTGC 48146

RESULT 39
US-10-756-149-2215/c
; Sequence 2215, Application US/10756149
; Publication No. US20050181375A1
; GENERAL INFORMATION:
; APPLICANT: Aziz, Natasha
; APPLICANT: Zlotnik, Albert
; TITLE OF INVENTION: NOVEL METHODS OF DIAGNOSIS OF METASTATIC CANCER, COMPOSITIONS AND
; TITLE OF INVENTION: METHODS OF SCREENING FOR MODULATORS OF METASTATIC CANCER
; FILE REFERENCE: file
```

db 2 GCGGTATATCCAGCACTTTGGGAGGCTTAAGGTGGGGGATACCTGAGGTCAAGAGTTC 61

QY 143 GAGACCATCTCGGCCAATGTTGAAAACCCGCTTTTACTAAAAATACAAAAATAGCTG 2022

db 242 CAGAGTGAGACTCCGCTCTCAAAAAACAAAAACAAAAACAAAAAC 301

```
QY 363 CA 364
DB 302 CA 303

RESULT 42
US-10-301-480-146767
; Sequence 146767, Application US/10301480
; Publication No. US20060057564A1
; GENERAL INFORMATION:
; APPLICANT: Wang, David G.
; TITLE OF INVENTION: Identification and Mapping of Single Nucleotide Polymorphisms
; FILE REFERENCE: 108827.137
; CURRENT APPLICATION NUMBER: US/10/301,480
; PRIOR FILING DATE: 2002-11-21
; PRIOR APPLICATION NUMBER: US 10/215,598
; PRIOR FILING DATE: 2002-08-09
; PRIOR APPLICATION NUMBER: US 60/311,695
; SOFTWARE: FastSeq for Windows Version 4.0
; NUMBER OF SEQ ID NOS: 1226818
; SEQ ID NO 146767
; LENGTH: 599
; TYPE: DNA
; ORGANISM: Homo sapien
US-10-301-480-146767

Query Match      37.9%; Score 151.6; DB 12; Length 599;
Best Local Similarity 74.2%; Pred. No. 4.2e-36;
Matches 224; Conservative 2; Mismatches 56; Indels 20; Gaps 2;

QY 83 GCCTGTATCCACAGCTTGGGAGGCCAAGTGGCGGATCAGTGAAGTCAAGATC 142
DB 2 GCCTGTATCCACAGCTTGGGAGGCCAAGTGGCGGATCAGTGAAGTCAAGATC 61

QY 143 GAGACCATCTGGGCAACATGATGTAACCCGCTCTTCTAAATAACAAAAATAGCTG 202
DB 62 GAGACCATCTGGGCAACATGATGTAACCCGCTCTTCTAAATAACAAAAATAGCTG 121

QY 203 GGCATGTGGCAGACACCTGTAGTCCCACTACTCAGAGCC---GGAGTTGCAATGA 258
DB 122 GACATGTGGCAGACACCTGTAGTCCCACTACTTGGAGGCTGAGCAATCATTGA 181

QY 259 GCTGAGATCGCAGAGTGGCCGAATTCACAGATCA-----CAGAGTGA 302
DB 182 ACCCGGAGGAGGAGTTGAGTGGCCAGATCAGCCACTGCACCTTCAAGCTTGA 241

QY 303 CAGAGTGAAGACCGCTCTCAAAAAACAACAACAAAAAACAATAGACATTGTC 362
DB 242 CAGAGTGAAGACCTCGCTCAAAAAACAACAACAAAAAACAATAGACAAAAAC 301

QY 363 CA 364
DB 302 CA 303

RESULT 43
US-10-301-480-760176
; Sequence 760176, Application US/10301480
; Publication No. US20060057564A1
; GENERAL INFORMATION:
; APPLICANT: Wang, David G.
; TITLE OF INVENTION: Identification and Mapping of Single Nucleotide Polymorphisms
; FILE REFERENCE: 108827.137
; CURRENT APPLICATION NUMBER: US/10/301,480
; PRIOR FILING DATE: 2002-11-21
; PRIOR APPLICATION NUMBER: US 10/215,598
; PRIOR FILING DATE: 2002-08-09
; PRIOR APPLICATION NUMBER: US 60/311,695
; SOFTWARE: FastSeq for Windows Version 4.0
; NUMBER OF SEQ ID NOS: 1226818
```

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; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 760176
; LENGTH: 599
; TYPE: DNA
; ORGANISM: Homo sapien
US-10-301-480-760176

Query Match      37.9%; Score 151.6; DB 12; Length 599;
Best Local Similarity 74.2%; Pred. No. 4.2e-36;
Matches 224; Conservative 2; Mismatches 56; Indels 20; Gaps 2;

QY 83 GCCTGTATCCACAGCTTGGGAGGCCAAGTGGCGGATCAGTGAAGTCAAGATC 142
DB 2 GCCTGTATCCACAGCTTGGGAGGCCAAGTGGCGGATCAGTGAAGTCAAGATC 61

QY 143 GAGACCATCTGGGCAACATGATGTAACCCGCTCTTCTAAATAACAAAAATAGCTG 202
DB 62 GAGACCATCTGGGCAACATGATGTAACCCGCTCTTCTAAATAACAAAAATAGCTG 121

QY 203 GGCATGTGGCAGACACCTGTAGTCCCACTACTCAGAGCC---GGAGTTGCAATGA 258
DB 122 GACATGTGGCAGACACCTGTAGTCCCACTACTTGGAGGCTGAGCAATCATTGA 181

QY 259 GCTGAGATCGCAGAGTGGCCGAATTCACAGATCA-----CAGAGTGA 302
DB 182 ACCCGGAGGAGGAGTTGAGTGGCCAGATCAGCCACTGCACCTTCAAGCTTGA 241

QY 303 CAGAGTGAAGACCGCTCTCAAAAAACAACAACAAAAAACAATAGACATTGTC 362
DB 242 CAGAGTGAAGACCTCGCTCAAAAAACAACAACAAAAAACAATAGACAAAAAC 301

QY 363 CA 364
DB 302 CA 303

RESULT 44
US-10-450-826-93/c
; Sequence 93, Application US/10450826
; Publication No. US20040101818A1
; GENERAL INFORMATION:
; APPLICANT: Mertz, Darren
; APPLICANT: Axelrod, Douglas W.
; APPLICANT: Cook, Jonathan S.
; APPLICANT: Eistein, Neelam
; APPLICANT: Houghton, Richard
; APPLICANT: Mertz, Lawrence
; TITLE OF INVENTION: Gene Expression Profiles Associated with Osteoblast Differentiation
; FILE REFERENCE: 044921-5039-WO
; CURRENT APPLICATION NUMBER: US/10/450,826
; CURRENT FILING DATE: 2003-06-18
; PRIOR APPLICATION NUMBER: US 60/255,882
; PRIOR FILING DATE: 2000-12-18
; PRIOR APPLICATION NUMBER: US 60/285,691
; PRIOR FILING DATE: 2001-04-24
; NUMBER OF SEQ ID NOS: 149
; SOFTWARE: Patentin Ver. 2.1
; SEQ ID NO 93
; LENGTH: 169739
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; OTHER INFORMATION: Genbank Accession No. AC005082
US-10-450-826-93

Query Match      37.9%; Score 151.4; DB 8; Length 169739;
Best Local Similarity 70.9%; Pred. No. 4.6e-35;
Matches 229; Conservative 1; Mismatches 87; Indels 6; Gaps 2;

QY 30 GGAACCAATATTAATAGACATTGTCAGGCCAGCATGACATGCTGAATGCTGTA 89
DB 125892 GGTAAAGAAATCTTAGTTAAAGATCTAAAGTTGGCCAGGCAAGTGGCTAACGCTGTA 125833
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? Publication No. US20050272054A1
? GENERAL INFORMATION:
? APPLICANT: CARGILL, Michele et al.
? TITLE OF INVENTION: GENETIC POLYMORPHISMS ASSOCIATED WITH
? TITLE OF INVENTION: CARDIOVASCULAR DISORDERS AND DRUG RESPONSE, METHODS OF
? TITLE OF INVENTION: DETECTION AND USES THEREOF
? FILE REFERENCE: CU001559
? CURRENT APPLICATION NUMBER: US/10/995,561
? CURRENT FILING DATE: 2004-11-24
? NUMBER OF SEQ ID NOS: 85702
? SOFTWARE: FastSeq for Windows Version 4.0
? SEQ ID NO 13489
? LENGTH: 195998
? TYPE: DNA
? ORGANISM: Homo sapiens
? FEATURE:
? NAME/KEY: misc_feature
? LOCATION: (1...195998)
? OTHER INFORMATION: n = A,T,C or G, or insertion/deletion polymorphism (see Tables 1
? S-10-995-561-13489

```

Query Match	37.8%;	Score 151;	DB 10;	Length 195998;
Best Local Similarity	76.8%;	Pred. No. 6.5e-35;		
Matches 199;	Conservative	1;	Mismatches 51;	Indels 8; Gaps 1;
QY	81	ATGCGCTGTATATCCGACGACCTTGGGAGGCGCAAGGTGGGCGGATCACTTGAGGTCAAGAGA	140	
Db	141321	ACGCCCTGTATATCCGACGACCTTGGGAAACCGAGAGGGGTGGGTCACTTGAGGTCAAGAGT	141262	
QY	141	TGAGAGCATCTCTGGACCAATGGGTGAAACCCCGCTCTTTACTAAAAATACAAAAAATAGC	200	
Db	141261	TGAGACCAAGCCTGGCCCAATGGTGAAGAACCCCGTCTCTATGAAAAGTACAAAACTAGC	141202	
QY	201	TGGCGATGGTGGCAACACTCTGTAGTCCAGCTACTCAGAGGCCGAGATTGCAGTGAGC	260	
Db	141201	CGGGCATGGTGGCGGGCCCTGTATATCCCAAGCTACTCAGAGCGCAGAGGTGTGACGTGAAC	141142	
QY	261	TGAGATTCGAGAGTGAAGCGGAAATCAAGATCAAGAGTGAAGCAGATGAGACCKCGTCT	320	
Db	141141	CGAAATTCG-----TGCCACTGACTCCAGGCTGGGTGACGTGACGAGACTCCGCTCT	141090	
QY	321	CAAAAAACAACAAAAA	339	
Db	141089	CAAAAAAAAAAAAAAAAAA	141071	

```

RESULT 47
US-09-764-860-797/c
; Sequence 797, Application US/09764860
; Patent No. US20020094953a1
GENERAL INFORMATION:
APPLICANT: Rosen et al.
TITLE OF INVENTION: Nucleic Acids, Proteins, and Antibodies
FILE REFERENCE: PC008
CURRENT APPLICATION NUMBER: US/09/764,860
CURRENT FILING DATE: 2001-01-17
Prior application data removed - consult PAMM or file wrapper
NUMBER OF SEQ ID NOS: 1198
SOFTWARE: PatentIn Ver. 2.0
SEQ ID NO 797
LENGTH: 32146
TYPE: DNA
ORGANISM: Homo sapiens
US-09-764-860-797

Query Match      37.7%; Score 150.6; DB 3; Length 32146;
Best Local Similarity 68.5%; Pred. No. 4.2e-35;
Matches 224; Conservative 1; Mismatches 95; Indels 7; Gaps 1;

      81 ATGCCGTATCCACACCTTCGGAGGCCAAGTGGCGCGATCACTGAGGTCAAGGA 140
      |||||
Db      2437 ACCCCGTATGTAGCACTTTGGAGGCCAAGCGGAGGAGTCACTTGAGGTCAAGAGT 2378

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PRIOR FILING DATE: 2000-08-22
PRIOR APPLICATION NUMBER: 60/225, 759
PRIOR FILING DATE: 2000-08-14
PRIOR APPLICATION NUMBER: 60/225, 213
PRIOR FILING DATE: 2000-08-14
PRIOR APPLICATION NUMBER: 60/227, 182
PRIOR FILING DATE: 2000-08-22
PRIOR APPLICATION NUMBER: 60/225, 214
PRIOR FILING DATE: 2000-08-14
PRIOR APPLICATION NUMBER: 60/235, 836
PRIOR FILING DATE: 2000-09-27
PRIOR APPLICATION NUMBER: 60/230, 438
PRIOR FILING DATE: 2000-09-06
PRIOR APPLICATION NUMBER: 60/215, 135
PRIOR FILING DATE: 2000-06-30
PRIOR APPLICATION NUMBER: 60/225, 266
PRIOR FILING DATE: 2000-08-14
PRIOR APPLICATION NUMBER: 60/249, 218
PRIOR FILING DATE: 2000-11-17
PRIOR APPLICATION NUMBER: 60/249, 208
PRIOR FILING DATE: 2000-11-17
PRIOR APPLICATION NUMBER: 60/249, 213
PRIOR FILING DATE: 2000-11-17
PRIOR APPLICATION NUMBER: 60/249, 212
PRIOR FILING DATE: 2000-11-17
PRIOR APPLICATION NUMBER: 60/249, 207
PRIOR FILING DATE: 2000-11-17
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PRIOR FILING DATE: 2000-11-08
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PRIOR FILING DATE: 2000-09-08

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DB 2317 AGGGCATGTGCGGCGCACCTGTAGTCCAGCTGCTTGGAGAGCTGAGCCAGAGAAATTG 2258
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QY 374 CCCAGCTATTGCGAGAGCAACAAAAG 400
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Sequence 797, Application US/10212872
Publication No. US20030215893A1
GENERAL INFORMATION:
APPLICANT: Rosen et al.
TITLE OF INVENTION: Nucleic Acids, Proteins, and Antibodies
FILE REFERENCE: PC008C2
CURRENT APPLICATION NUMBER: US/10/212,872
CURRENT FILING DATE: 2002-08-07
Prior application removed - See File Wrapper or Palm
NUMBER OF SEQ ID NOS: 1198
SOFTWARE: PatentIn Ver. 2.0
SEQ ID NO 797
LENGTH: 32146
TYPE: DNA
ORGANISM: Homo sapiens
US-10-212-872-797

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DB 2437 AGCGCTGTAAATGCTAGCACTTTGGAGGCGCAAGGCGGAGATCACTTGAGGTCAAGGT 2378
QY 141 TCGAGACCATCTGCGCCCAACATGTTGAACCCCGCTTTACTTAAATAACAAAATATGAC 200
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RESULT 50

US-09-893-348-9/c
Sequence 9, Application US/09893348
Patent No. US20020072493A1
GENERAL INFORMATION:
APPLICANT: EISENBACH-SCHWARTZ, Michael
APPLICANT: COHEN, Irvin R.
APPLICANT: BESERMAN, Plette
APPLICANT: MOSONEGO, Alon
APPLICANT: MOLEEM, Gila
TITLE OF INVENTION: ACTIVATED T-CELLS, NERVOUS SYSTEM-SPECIFIC ANTIGENS AND THEIR USE
FILE REFERENCE: EIS-SCHWARTZ-2A
CURRENT APPLICATION NUMBER: US/09/893,348
PRIOR FILING DATE: 2001-06-28
PRIOR APPLICATION NUMBER: US 09/314,161
PRIOR FILING DATE: 1999-05-19
PRIOR APPLICATION NUMBER: US 09/218,277
PRIOR FILING DATE: 1998-12-22
PRIOR APPLICATION NUMBER: PCT/US98/14715
PRIOR FILING DATE: 1998-07-21
PRIOR APPLICATION NUMBER: IL 124500
PRIOR FILING DATE: 1998-05-19
NUMBER OF SEQ ID NOS: 29
SOFTWARE: PatentIn version 3.1
SEQ ID NO 9
LENGTH: 17538
TYPE: DNA
ORGANISM: Homo sapiens
US-09-893-348-9

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Job time : 1466 secs

GenCore version 5.1.9
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On nucleic - nucleic search, using sw model

Run on: July 17, 2006, 21:21:05 ; Search time 407 Seconds

(without alignments)
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Total number of hits satisfying chosen parameters: 1772710

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Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

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ALIGNMENTS

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; Publication No. US20060134663A1
; GENERAL INFORMATION:
; APPLICANT: Hairkin, Paul
; APPLICANT: Mulligan, Patrick
; TITLE OF INVENTION: Transcriptome Microarray Technology and
; FILE REFERENCE: 55815-0102 (319189)
; CURRENT FILING DATE: 2005-11-03
; PRIOR APPLICATION NUMBER: EP 04105479.2
; PRIOR FILING DATE: 2004-11-03

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; PRIOR FILING DATE: 2004-11-03
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; PRIOR FILING DATE: 2004-11-03
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; PRIOR FILING DATE: 2004-11-03
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; PRIOR FILING DATE: 2004-11-03
; PRIOR APPLICATION NUMBER: EP 04105484.2
; PRIOR FILING DATE: 2004-11-03
; PRIOR APPLICATION NUMBER: US 60/700,293
; NUMBER OF SEQ ID NOS: 483996
; SOFTWARE: PatentIn version 3.3
; SEQ ID NO 23004
; LENGTH: 216387
; TYPE: DNA
; ORGANISM: Homo Sapiens
US-11-266-748A-23004
Query Match 39.0%; Score 155.8; DB 8; Length 216387;
Best Local Similarity 77.0%; Pred. No. 9.5e-24;
Matches 218; Conservative 1; Mismatches 53; Indels 11; Gaps 2;
QY 81 ATGCTGTAATCCAGACCTCGGAGGCGCAAGGTGGGCGGATCACCCTGAGTCAAGAGA 140
DB 195538 ATGCTGTAATCCAGACCTCGGAGGCGCAAGGTGGGCGGATCACCCTGAGTCAAGAGA 195597
QY 141 TCGAGACCATCTGCGCCCAACATGCTGAAACCCCGCTTTACTAAATATCAAAAATATGC 200
DB 195598 TCGAGACCATCTGCGCCCAACATGCTGAAACCCCGCTTTACTAAATATCAAAAATATGC 195657
QY 201 TGGGCACTGTCGACACACCTGTGTCTCCAGCTACTAGAGA-----GCCGAGATTGC 253
DB 195658 TGGGCGTGTGTGTATGATCTGTGTATGCTCCAGCTACTCGGAGGCTGAGGCAAGAGATTG 195717
QY 254 AGTAGCTGATATGCGAGTAGGAGCCGGAATCACAGAT---CACAGATGAGCAGAGTGC 309
DB 195718 CTGGAACCTGAGAGCGACAGGCTGATGTAACCAAGATATCTCCAGCTTAGTGAACAGCG 195777
QY 310 AGAGCCCTCTCAAAAACAAACAAACAAACAAACAAACAAACAAACAAACAAACAAAC 352
DB 195778 AGAGCCCTCTCAAAAACAAACAAACAAACAAACAAACAAACAAACAAACAAACCTTA 195820

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RESULT 2
US-11-266-748A-204137/c
; Sequence 204137, Application US/11266748A
; Publication No. US20060134663A1
; GENERAL INFORMATION:
; APPLICANT: Hairkin, Paul
; APPLICANT: Mulligan, Karl
; TITLE OF INVENTION: Transcriptome Microarray Technology and
; FILE REFERENCE: 55815-0102 (319189)
; CURRENT FILING DATE: 2005-11-03
; PRIOR APPLICATION NUMBER: EP 04105479.2
; PRIOR FILING DATE: 2004-11-03
; PRIOR APPLICATION NUMBER: EP 04105482.6
; PRIOR FILING DATE: 2004-11-03
; PRIOR APPLICATION NUMBER: EP 04105483.4
; PRIOR FILING DATE: 2004-11-03
; PRIOR APPLICATION NUMBER: EP 04105507.0
; PRIOR FILING DATE: 2004-11-03
; PRIOR APPLICATION NUMBER: EP 04105485.9
; PRIOR FILING DATE: 2004-11-03
; PRIOR APPLICATION NUMBER: EP 04105484.2
; PRIOR FILING DATE: 2004-11-03
; PRIOR APPLICATION NUMBER: US 60/662,276

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PRIOR FILING DATE: 2005-03-14
PRIOR APPLICATION NUMBER: US 60/700,293
PRIOR FILING DATE: 2005-07-18
NUMBER OF SEQ ID NOS: 483996
SOFTWARE: PatentIn version 3.3
SEQ ID NO 204137
LENGTH: 1000
TYPE: DNA
ORGANISM: Homo Sapiens
US-11-266-748A-204137

Query Match 37.8%; Score 151; DB 8; Length 1000;
Best Local Similarity 74.4%; Pred. No. 7,2e-23;
Matches 206; Conservative 1; Mismatches 61; Indels 9; Gaps 1;

QY 81 ATGCTGTATATCCAGCACTTGGGAGGCGCAAGTGGCGGATCACTGAGGTCAAGAGA 140
DB 303 ATGCTGTATATCCAGCACTTGGGAGGCGTGAAGTCATCACTTAAGTCAAGAGT 244
QY 141 TCGAGACCATCTGGGCAACATGAGTGAACCCCGTCTTAAATAATCAAAAAATAGC 200
DB 243 TCAAGACCAAGCTGGGCAACATGAGTGAACCTGTCTGTCTTAAATAATCAAAAAATAGC 184
QY 201 TGGGATGATGGGCAACACCTGTAGTCCAGCTACTCAGAGCCGAGATTCAGATGAGC 260
DB 183 CAGGATAGTGGGCAAGCACTGTATATCCAGCTACTTGGAGCCGAGGTTCAGATGAGC 124
QY 261 TGAAGTCGAGAGTGAGCCGAAATCAAGATCAAGAGTGAAGCAGATGAGCCKCGTCT 320
DB 123 TGAGTTCGACCACTGTCTTAAAGCTGGGTGA-----CAGAGCAAGACTGTCT 73
QY 321 CAAAAACAACAACAAAAACCAAAACCAATAGCA 357
DB 72 CAAAAACAAAAACAAAAACCAAAACCAAAACA 36

RESULT 3

US-11-266-748A-391815
Sequence 391815, Application US/11266748A
Publication No. US20060134663A1
GENERAL INFORMATION:
APPLICANT: Harkin, Paul
APPLICANT: Johnston, Patrick
TITLE OF INVENTION: Transcription Microarray Technology and
TITLE OF INVENTION: Methods of Using the Same
FILE REFERENCE: 55815-0102 (319189)
CURRENT APPLICATION NUMBER: US/11/266,748A
PRIOR FILING DATE: 2005-11-03
PRIOR APPLICATION NUMBER: EP 04105479.2
PRIOR FILING DATE: 2004-11-03
PRIOR APPLICATION NUMBER: EP 04105482.6
PRIOR FILING DATE: 2004-11-03
PRIOR APPLICATION NUMBER: EP 04105483.4
PRIOR FILING DATE: 2004-11-03
PRIOR APPLICATION NUMBER: EP 04105507.0
PRIOR FILING DATE: 2004-11-03
PRIOR APPLICATION NUMBER: EP 04105485.9
PRIOR FILING DATE: 2004-11-03
PRIOR APPLICATION NUMBER: EP 04105484.2
PRIOR FILING DATE: 2004-11-03
PRIOR APPLICATION NUMBER: US 60/662,276
PRIOR FILING DATE: 2005-03-14
PRIOR APPLICATION NUMBER: US 60/700,293
NUMBER OF SEQ ID NOS: 483996
SOFTWARE: PatentIn version 3.3
SEQ ID NO 391815
LENGTH: 1000
TYPE: DNA
ORGANISM: Homo Sapiens
US-11-266-748A-391815

Query Match 37.4%; Score 149.4; DB 8; Length 1000;
Best Local Similarity 74.0%; Pred. No. 1.6e-22;
Matches 205; Conservative 1; Mismatches 62; Indels 9; Gaps 1;

QY 81 ATGCTGTATATCCAGCACTTGGGAGGCGCAAGTGGCGGATCACTGAGGTCAAGAGA 140
DB 732 AGGCTGTATATCCAGCACTTGGGAGGCGTCAAGTGGGCAATCACTTGAATCAGAGT 791
QY 141 TCGAGACCATCTGGGCAACATGAGTGAACCCCGTCTTAAATAATCAAAAAATAGC 200
DB 792 TCAAGACCAAGCTGGGCAACATGAGTGAACCCCGTCTTAAATAATCAAAAAATAGC 851
QY 201 TGGGATGATGGGCAACACCTGTAGTCCAGCTACTCAGAGCCGAGATTCAGATGAGC 260
DB 852 CATGATGATGGGCAACATGAGTGAACCACTTAAAGGAGGCAAGATTCAGATGAGC 911
QY 261 TGAAGTCGAGAGTGAGCCGAAATCAAGATCAAGAGTGAAGCAGATGAGCCKCGTCT 320
DB 912 TGAGTTCGACCACTGTCTTAAAGCTGGGTGA-----GCTGGGCTGGCAAGCAGATCTCATCT 962
QY 321 CAAAAACAACAACAAAAACCAAAACCAATAGCA 357
DB 963 CAAAAACAAAAACAAAAACCAAAACCAAAACA 999

RESULT 4

US-11-266-748A-482533/c
Sequence 482533, Application US/11266748A
Publication No. US20060134663A1
GENERAL INFORMATION:
APPLICANT: Harkin, Paul
APPLICANT: Johnston, Patrick
TITLE OF INVENTION: Transcription Microarray Technology and
TITLE OF INVENTION: Methods of Using the Same
FILE REFERENCE: 55815-0102 (319189)
CURRENT APPLICATION NUMBER: US/11/266,748A
PRIOR FILING DATE: 2005-11-03
PRIOR APPLICATION NUMBER: EP 04105479.2
PRIOR FILING DATE: 2004-11-03
PRIOR APPLICATION NUMBER: EP 04105482.6
PRIOR FILING DATE: 2004-11-03
PRIOR APPLICATION NUMBER: EP 04105483.4
PRIOR FILING DATE: 2004-11-03
PRIOR APPLICATION NUMBER: EP 04105507.0
PRIOR FILING DATE: 2004-11-03
PRIOR APPLICATION NUMBER: EP 04105485.9
PRIOR FILING DATE: 2004-11-03
PRIOR APPLICATION NUMBER: EP 04105484.2
PRIOR FILING DATE: 2004-11-03
PRIOR APPLICATION NUMBER: US 60/662,276
PRIOR FILING DATE: 2005-03-14
PRIOR APPLICATION NUMBER: US 60/700,293
PRIOR FILING DATE: 2005-07-18
NUMBER OF SEQ ID NOS: 483996
SOFTWARE: PatentIn version 3.3
SEQ ID NO 482533
LENGTH: 1000
TYPE: DNA
ORGANISM: Homo Sapiens
US-11-266-748A-482533

Query Match 37.4%; Score 149.4; DB 8; Length 1000;
Best Local Similarity 74.0%; Pred. No. 1.6e-22;
Matches 205; Conservative 1; Mismatches 62; Indels 9; Gaps 1;

QY 81 ATGCTGTATATCCAGCACTTGGGAGGCGCAAGTGGCGGATCACTGAGGTCAAGAGA 140
DB 269 AGGCTGTATATCCAGCACTTGGGAGGCGTCAAGTGGGCAATCACTTGAATCAGAGT 210
QY 141 TCGAGACCATCTGGGCAACATGAGTGAACCCCGTCTTAAATAATCAAAAAATAGC 200
DB 209 TCAAGACCAAGCTGGGCAACATGAGTGAACCCCGTCTTAAATAATCAAAAAATAGC 150


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RESULT 7
US-11-266-748A-403127
; Sequence 403127, Application US/11266748A
; Publication No. US2006013463A1
; GENERAL INFORMATION:
; APPLICANT: Harkin, Paul
; APPLICANT: Johnston, Patrick
; APPLICANT: Mulligan, Karl
; TITLE OF INVENTION: Transcription Microarray Technology and
; FILE REFERENCE: 55815-0102 (319189)
; CURRENT FILING DATE: 2005-11-03
; PRIOR FILING DATE: 2004-11-03
; PRIOR APPLICATION NUMBER: EP 04105479.2
; PRIOR FILING DATE: 2004-11-03
; PRIOR APPLICATION NUMBER: EP 04105482.6
; PRIOR FILING DATE: 2004-11-03
; PRIOR APPLICATION NUMBER: EP 04105483.4
; PRIOR FILING DATE: 2004-11-03
; PRIOR APPLICATION NUMBER: EP 04105507.0
; PRIOR FILING DATE: 2004-11-03
; PRIOR APPLICATION NUMBER: EP 04105485.9
; PRIOR FILING DATE: 2004-11-03
; PRIOR APPLICATION NUMBER: EP 04105484.2
; PRIOR FILING DATE: 2004-11-03
; PRIOR APPLICATION NUMBER: US 60/662,276
; PRIOR FILING DATE: 2005-03-14
; PRIOR APPLICATION NUMBER: US 60/700,293
; PRIOR FILING DATE: 2005-07-18
; NUMBER OF SEQ ID NOS: 483996
; SOFTWARE: PatentIn version 3.3
; SEQ ID NO 403127
; LENGTH: 1000
; TYPE: DNA
; ORGANISM: Homo Sapiens
US-11-266-748A-403127

Query Match      37.0%; Score 147.8; DB 8; Length 1000;
Best Local Similarity 68.4%; Pred. No. 3.4e-22;
Matches 221; Conservative 1; Mismatches 93; Indels 8; Gaps 1;

QY 33 ACCCAATATTAAATAGACATTTGTGAGGCCAGGATGACACTGCTGATGCTGTATATC 92
DB 677 ACACATTTTACATTAAATAATACGTGTCAGGCCATGCTGTGCTGACGCTTATATC 736
QY 93 CCAGCACTTCGGAGGCCAGAGTGGCGGATCACCTGAGTCAAGATCGAGACCATCC 152
DB 737 CCAGCACTTTGGAGGCCAGAGTGGCGGATCACCGGAGTCAAGAGTTCCAGACCAACC 796
QY 153 TGGCAACATGTTGTAACCCCGTCTTTACTTAAATAATCAAAAAATAGCTGGGCAATGG 212
DB 797 TTGCAACATGTTGTAACCCCGTCTTTACTTAAATAATCAAAAAATAGCTGGGCAATGG 856
QY 213 CACACACCTGTAGTCCAGCTACTCAGAG-----CCGAGATTGCACTGAGCTGAG 264
DB 857 CAGGCACTGTATATCCAGTACTAGGAGGCTTTGAACCCAGAGGCAAGAGTTGCGAG 916
QY 265 ATGCAAGTGAAGCCGAATCAAGATCAAGATGAGAGTGAAGAGTGAAGCCKCGTCTCAA 324
DB 917 CGAGCTGAGATCGCGCCACTGCACTCCAGCTGGGTATAGATGATTCAGTCTCAA 976
QY 325 AACCAACAACAAAAACAAAAA 347
DB 977 AAAAAAAAAAAAAAAAAAAAAA 999

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; APPLICANT: Mulligan, Karl
; TITLE OF INVENTION: Transcription Microarray Technology and
; FILE REFERENCE: 55815-0102 (319189)
; CURRENT FILING DATE: 2005-11-03
; PRIOR FILING DATE: 2004-11-03
; PRIOR APPLICATION NUMBER: EP 04105479.2
; PRIOR FILING DATE: 2004-11-03
; PRIOR APPLICATION NUMBER: EP 04105482.6
; PRIOR FILING DATE: 2004-11-03
; PRIOR APPLICATION NUMBER: EP 04105483.4
; PRIOR FILING DATE: 2004-11-03
; PRIOR APPLICATION NUMBER: EP 04105507.0
; PRIOR FILING DATE: 2004-11-03
; PRIOR APPLICATION NUMBER: EP 04105485.9
; PRIOR FILING DATE: 2004-11-03
; PRIOR APPLICATION NUMBER: EP 04105484.2
; PRIOR FILING DATE: 2004-11-03
; PRIOR APPLICATION NUMBER: US 60/662,276
; PRIOR FILING DATE: 2005-03-14
; PRIOR APPLICATION NUMBER: US 60/700,293
; PRIOR FILING DATE: 2005-07-18
; NUMBER OF SEQ ID NOS: 483996
; SOFTWARE: PatentIn version 3.3
; SEQ ID NO 474173
; LENGTH: 1000
; TYPE: DNA
; ORGANISM: Homo Sapiens
US-11-266-748A-474173

Query Match      37.0%; Score 147.8; DB 8; Length 1000;
Best Local Similarity 68.4%; Pred. No. 3.4e-22;
Matches 221; Conservative 1; Mismatches 93; Indels 8; Gaps 1;

QY 33 ACCCAATATTAAATAGACATTTGTGAGGCCAGGATGACACTGCTGATGCTGTATATC 92
DB 324 ACACATTTTACATTAAATAATACGTGTCAGGCCATGCTGTGCTGACGCTTATATC 265
QY 93 CCAGCACTTCGGAGGCCAGAGTGGCGGATCACCTGAGTCAAGATCGAGACCATCC 152
DB 264 CCAGCACTTTGGAGGCCAGAGTGGCGGATCACCGGAGTCAAGAGTTCCAGACCAACC 205
QY 153 TGGCAACATGTTGTAACCCCGTCTTTACTTAAATAATCAAAAAATAGCTGGGCAATGG 212
DB 204 TTGCAACATGTTGTAACCCCGTCTTTACTTAAATAATCAAAAAATAGCTGGGCAATGG 145
QY 213 CACACACCTGTAGTCCAGCTACTCAGAG-----CCGAGATTGCACTGAGCTGAG 264
DB 144 CAGGCACTGTATATCCAGTACTAGGAGGCTTTGAACCCAGAGGCAAGAGTTGCGAG 85
QY 265 ATGCAAGTGAAGCCGAATCAAGATCAAGATGAGAGTGAAGAGTGAAGCCKCGTCTCAA 324
DB 84 CGAGCTGAGATCGCGCCACTGCACTCCAGCTGGGTATAGATGATTCAGTCTCAA 25
QY 325 AACCAACAACAAAAACAAAAA 347
DB 24 AAAAAAAAAAAAAAAAAAAAAA 2

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RESULT 8
US-11-266-748A-474173/C
; Sequence 474173, Application US/11266748A
; Publication No. US2006013463A1
; GENERAL INFORMATION:
; APPLICANT: Harkin, Paul
; APPLICANT: Johnston, Patrick

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RESULT 9
US-11-319-952-56
; Sequence 56, Application US/11319952
; Publication No. US2006013414A1
; GENERAL INFORMATION:
; APPLICANT: Yousef, George M.
; APPLICANT: Diamandis, Efstherios
; TITLE OF INVENTION: Novel Human Kallikrein-like Genes
; FILE REFERENCE: WTS3USA
; CURRENT FILING DATE: 2005-12-28
; PRIOR FILING DATE: 2001-09-10
; PRIOR APPLICATION NUMBER: US/09/936,271
; PRIOR APPLICATION NUMBER: PCT/CA00/00258

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; PRIOR FILING DATE: 2000-03-09
; PRIOR APPLICATION NUMBER: US 60/124,260
; PRIOR FILING DATE: 1999-03-11
; PRIOR APPLICATION NUMBER: US 60/127,386
; PRIOR FILING DATE: 1999-04-01
; PRIOR APPLICATION NUMBER: US 60/144,919
; PRIOR FILING DATE: 1999-07-21
; NUMBER OF SEQ ID NOS: 97
; SOFTWARE: PatentIn version 3.2
; SEQ ID NO 56
; LENGTH: 11820
; TYPE: DNA
; ORGANISM: Homo sapiens
US-11-319-952-56

Query Match          36.8%; Score 147.6; DB 8; Length 11820;
Best Local Similarity 75.4%; Pred. No. 4.2e-22;
Matches 199; Conservative 1; Mismatches 55; Indels 9; Gaps 1;

QY 84 CCTGTAATCCAGCATTTCGGGAGGCGCAAGTGGCGGATCACTGAGGTCAAGAGATCG 143
DB 2706 CCTGTAATCCCGCGCATTTGGGAAAGCCAAAGGCAAGGCAAGTTGCTTGAGGCCAGAGTTCA 2765
QY 144 AGACCATCTGCGCCCAACATGGGAAACCCCGCTTTACTTAAATAACAAAAATATAGCTGG 203
DB 2766 AGACCATCTGCGCCCAACATGGGAAACCCCGCTTTACTTAAATAACAAAAATATAGCTGG 2825
QY 204 GCATGTGGGACACACCTGTAGTCCCACTACTCAGAGCCGAGATTGCAAGTGAAGCTGA 263
DB 2826 ACATGTGGGACACAGTGTGCTGTATCTCCAGTACTCAAGAGGTGAGAGTTGCAAGTGAAGCTGA 2885
QY 264 GATGCGAAGTGAAGCCGAAATCAAGATCAAGAGTGAAGAGAGTGAACCCGCTCTAA 323
DB 2886 GATGCGAAGTGAAGCCGCTGCACTCAGCTGGGAGA-----CAGAGGAGACTTCATCTCAA 2936
QY 324 AAACACACACAAAAAACAACAAAAA 347
DB 2937 AACGAAAAACAAAAACAGCACAA 2960

RESULT 10
US-11-266-748A-219455
; Sequence 219455, Application US/11266748A
; Publication No. US20060134663A1
; GENERAL INFORMATION:
; APPLICANT: Harkin, Paul
; APPLICANT: Johnston, Patrick
; TITLE OF INVENTION: Transcriptome Microarray Technology and
; FILE REFERENCE: 55815-0102 (319189)
; CURRENT FILING DATE: US/11/266,748A
; PRIOR FILING DATE: 2005-11-03
; PRIOR APPLICATION NUMBER: EP 04105479.2
; PRIOR FILING DATE: 2004-11-03
; PRIOR APPLICATION NUMBER: EP 04105482.6
; PRIOR FILING DATE: 2004-11-03
; PRIOR APPLICATION NUMBER: EP 04105483.4
; PRIOR FILING DATE: 2004-11-03
; PRIOR APPLICATION NUMBER: EP 04105507.0
; PRIOR FILING DATE: 2004-11-03
; PRIOR APPLICATION NUMBER: EP 04105485.9
; PRIOR FILING DATE: 2004-11-03
; PRIOR APPLICATION NUMBER: EP 04105484.2
; PRIOR FILING DATE: 2004-11-03
; PRIOR APPLICATION NUMBER: US 60/662,276
; PRIOR FILING DATE: 2005-03-14
; PRIOR APPLICATION NUMBER: US 60/700,293
; PRIOR FILING DATE: 2005-07-18
; NUMBER OF SEQ ID NOS: 483996
; SOFTWARE: PatentIn version 3.3
; SEQ ID NO 219455
; LENGTH: 897
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; TYPE: DNA
; ORGANISM: Homo sapiens
US-11-266-748A-219455

Query Match          36.8%; Score 147.2; DB 8; Length 897;
Best Local Similarity 75.4%; Pred. No. 4.4e-22;
Matches 196; Conservative 1; Mismatches 59; Indels 4; Gaps 1;

QY 81 ATGCTGTAATCCAGCATTTCGGGAGGCGCAAGTGGCGGATCACTGAGGTCAAGAGA 140
DB 593 AGCGCTGTAATCTTGAAGACTTTGGGAGCGGAGGAGTGAATCACTGAGGTCAAGAGT 652
QY 141 TCGAGACCATCTGCGCCAAATGAGTGAACCCCGCTTTACTTAAATAACAAAAATATAGC 200
DB 653 TCGAGACCATCTGCGCCAAATGAGTGAACCCCGCTTTACTTAAATAACAAAAATATAGC 712
QY 201 TGGGCAATGTTGGACACACCTGTAGTCCAGTACTCAGAGAGCCGAGATTGCAAGTGAAGC 260
DB 713 TGGGTATGTGTGTCATGCTGTATCCAGCTACTAGGAGGCTGAGGCGAGAGAAATCA 772
QY 261 TGAATGCGACAGTGAAGCCGAAATCAAGATCAAGAGTGAAGAGTGAAGAGACCGCTCT 320
DB 773 CTGGAACCCAGAGTGAAGGCGACAGTGAAC---AGGCGCAGAGTGAAGACTCTGTCT 828
QY 321 CAAAAACACACAAAAAAC 340
DB 829 CAAAAACAAAAAAGAAAAAC 848

RESULT 11
US-11-266-748A-239441/c
; Sequence 239441, Application US/11266748A
; Publication No. US20060134663A1
; GENERAL INFORMATION:
; APPLICANT: Harkin, Paul
; APPLICANT: Johnston, Patrick
; TITLE OF INVENTION: Transcriptome Microarray Technology and
; FILE REFERENCE: 55815-0102 (319189)
; CURRENT FILING DATE: US/11/266,748A
; PRIOR FILING DATE: 2005-11-03
; PRIOR APPLICATION NUMBER: EP 04105479.2
; PRIOR FILING DATE: 2004-11-03
; PRIOR APPLICATION NUMBER: EP 04105482.6
; PRIOR FILING DATE: 2004-11-03
; PRIOR APPLICATION NUMBER: EP 04105483.4
; PRIOR FILING DATE: 2004-11-03
; PRIOR APPLICATION NUMBER: EP 04105507.0
; PRIOR FILING DATE: 2004-11-03
; PRIOR APPLICATION NUMBER: EP 04105485.9
; PRIOR FILING DATE: 2004-11-03
; PRIOR APPLICATION NUMBER: EP 04105484.2
; PRIOR FILING DATE: 2004-11-03
; PRIOR APPLICATION NUMBER: US 60/662,276
; PRIOR FILING DATE: 2005-03-14
; PRIOR APPLICATION NUMBER: US 60/700,293
; PRIOR FILING DATE: 2005-07-18
; NUMBER OF SEQ ID NOS: 483996
; SOFTWARE: PatentIn version 3.3
; SEQ ID NO 239441
; LENGTH: 897
; TYPE: DNA
; ORGANISM: Homo sapiens
US-11-266-748A-239441

Query Match          36.8%; Score 147.2; DB 8; Length 897;
Best Local Similarity 75.4%; Pred. No. 4.4e-22;
Matches 196; Conservative 1; Mismatches 59; Indels 4; Gaps 1;

QY 81 ATGCTGTAATCCAGCATTTCGGGAGGCGCAAGTGGCGGATCACTGAGGTCAAGAGA 140
DB 305 AGCGCTGTAATCTTGAAGACTTTGGGAGCGGAGGAGTGAATCACTGAGGTCAAGAGT 246
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141 TCGAGACCATCTGGCCAACTGTGAAACCCCGTCTTACTTAAATAACAAAAATAGC 200
245 TCGAGACCATCTGGCCAACTGTGAAACCCCGTCTTACTTAAATAACAAAAATAGC 186
201 TGGGATGTTGGGACACACCTGTAGTCCAGCTACTCAGAGACCGGAGATTGCAGTAGC 260
185 TGGGATGTTGGGACACACCTGTAGTCCAGCTACTCAGAGACCGGAGATTGCAGTAGC 126
261 TGAATTCGACAGTGAACCGGAAATACAGATACAGATGAGCAGAGTAGACACCCGCTCT 320
125 CTGTAACCCAGAGGAGGAGGACAGAGTACC-----AGGCGCCAGAGTAGAGCTGTCT 70
321 CAAAAACAACACAAAAAAC 340
69 CAAAAAGAAAAAGAAAAAC 50

RESULT 12
US-11-293-697-1113/c
; Sequence 1113, Application US/11293697
; Publication No. US20060105376A1
; GENERAL INFORMATION:
; APPLICANT: HELIX RESEARCH INSTITUTE
; TITLE OF INVENTION: Novel full length cDNA
; FILE REFERENCE: H1-A0106
; CURRENT APPLICATION NUMBER: US/11/293,697
; CURRENT FILING DATE: 2005-12-05
; PRIOR APPLICATION NUMBER: US/10/108,260
; PRIOR FILING DATE: 2002-03-28
; NUMBER OF SEQ ID NOS: 5458
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 1113
; LENGTH: 2909
; TYPE: DNA
; ORGANISM: Homo sapiens
US-11-293-697-1113

Query Match      36.8%; Score 147; DB 8; Length 2909;
Best Local Similarity 71.9%; Pred. No. 5.2e-22;
Matches 192; Conservative 0; Mismatches 75; Indels 0; Gaps 0;

141 ATGCGTGAATCCGACGACTTCGGAGGCGCAAGTGGGGATTCACCTGAGTCAAGAGA 140
678 ACGCTTAATATCCGACGACTTCGGAGGCGTGGAGCGGTGATCACTGAGGTGAGAGT 619
141 TCGAGACCATCTGGCCAACTGTGAAACCCCGTCTTACTTAAATAACAAAAATAGC 200
618 TCAAGACCATCTGGCCAACTGTGAAACCCCGTCTTACTTAAATAACAAAAATAGC 559
201 TGGGATGTTGGGACACACCTGTAGTCCAGCTACTCAGAGACCGGAGATTGCAGTAGC 260
558 TGGGATGTTGGGACACACCTGTAGTCCAGCTACTCAGAGACCGGAGATTGCAGTAGC 499
261 TGAATTCGACAGTGAACCGGAAATACAGATACAGATGAGCAGAGTAGACACCCGCTCT 320
498 CAAAGTTCACACAACTGCACTCCAGCTGGGTGACAGAGCAAGATCCGCTCTCCAAAAA 439
321 CAAAAACAACACAAAAAAC 347
438 TAAATTAATTAATTAATTAATTAATCA 412

RESULT 13
US-11-266-748A-23290/c
; Sequence 23290, Application US/11266748A
; Publication No. US20060134663A1
; GENERAL INFORMATION:
; APPLICANT: Harkin, Paul
; APPLICANT: Johnston, Patrick
; APPLICANT: Mulligan, Karl
; TITLE OF INVENTION: Transcription Microarray Technology and
; TITLE OF INVENTION: Methods of Using the Same
```

```
FILE REFERENCE: 55815-0102 (319189)
; CURRENT APPLICATION NUMBER: US/11/266,748A
; CURRENT FILING DATE: 2005-11-03
; PRIOR APPLICATION NUMBER: EP 04105479.2
; PRIOR FILING DATE: 2004-11-03
; PRIOR APPLICATION NUMBER: EP 04105482.6
; PRIOR FILING DATE: 2004-11-03
; PRIOR APPLICATION NUMBER: EP 04105483.4
; PRIOR FILING DATE: 2004-11-03
; PRIOR APPLICATION NUMBER: EP 04105507.0
; PRIOR FILING DATE: 2004-11-03
; PRIOR APPLICATION NUMBER: EP 04105485.9
; PRIOR FILING DATE: 2004-11-03
; PRIOR APPLICATION NUMBER: EP 04105484.2
; PRIOR FILING DATE: 2004-11-03
; PRIOR APPLICATION NUMBER: US 60/662,276
; PRIOR FILING DATE: 2005-03-14
; PRIOR APPLICATION NUMBER: US 60/700,293
; PRIOR FILING DATE: 2005-07-18
; NUMBER OF SEQ ID NOS: 48396
; SOFTWARE: PatentIn version 3.3
; SEQ ID NO 23290
; LENGTH: 112595
; TYPE: DNA
; ORGANISM: Homo Sapiens
US-11-266-748A-23290

Query Match      36.6%; Score 146.2; DB 8; Length 112595;
Best Local Similarity 78.5%; Pred. No. 9.1e-22;
Matches 175; Conservative 0; Mismatches 48; Indels 0; Gaps 0;

141 ATGCGTGAATCCGACGACTTCGGAGGCGCAAGTGGGGATTCACCTGAGTCAAGAGA 140
23489 ACGCTTAATATCCGACGACTTCGGAGGCGGAGGTGAGTCACTGAGGCCGAGAGT 23430
141 TCGAGACCATCTGGCCAACTGTGAAACCCCGTCTTACTTAAATAACAAAAATAGC 200
23429 TCGAGACCATCTGGCCAACTGTGAAACCCCGTCTTACTTAAATAACAAAAATAGC 23370
201 TGGGATGTTGGGACACACCTGTAGTCCAGCTACTCAGAGACCGGAGATTGCAGTAGC 260
23369 CCGGATGTTGGGACACACCTGTAGTCCAGCTACTCAGAGACCGGAGATTGCAGTAGC 23310
261 TGAATTCGACAGTGAACCGGAAATACAGATACAGATGAGTAGC 303
23309 TTTGAACCGGGGTGCGAAGGTTCGATGAGACCGGAGATTGCGC 23267

RESULT 14
US-11-266-748A-30503/c
; Sequence 30503, Application US/11266748A
; Publication No. US20060134663A1
; GENERAL INFORMATION:
; APPLICANT: Harkin, Paul
; APPLICANT: Johnston, Patrick
; APPLICANT: Mulligan, Karl
; TITLE OF INVENTION: Transcription Microarray Technology and
; TITLE OF INVENTION: Methods of Using the Same
FILE REFERENCE: 55815-0102 (319189)
; CURRENT APPLICATION NUMBER: US/11/266,748A
; CURRENT FILING DATE: 2005-11-03
; PRIOR APPLICATION NUMBER: EP 04105479.2
; PRIOR FILING DATE: 2004-11-03
; PRIOR APPLICATION NUMBER: EP 04105482.6
; PRIOR FILING DATE: 2004-11-03
; PRIOR APPLICATION NUMBER: EP 04105483.4
; PRIOR FILING DATE: 2004-11-03
; PRIOR APPLICATION NUMBER: EP 04105507.0
; PRIOR FILING DATE: 2004-11-03
; PRIOR APPLICATION NUMBER: EP 04105485.9
; PRIOR FILING DATE: 2004-11-03
; PRIOR APPLICATION NUMBER: EP 04105484.2
; PRIOR FILING DATE: 2004-11-03
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PRIOR APPLICATION NUMBER: US 60/662,276
PRIOR FILING DATE: 2005-03-14
PRIOR APPLICATION NUMBER: US 60/700,293
PRIOR FILING DATE: 2005-07-18
NUMBER OF SEQ ID NOS: 483996
SOFTWARE: PatentIn version 3.3
SEQ ID NO 30503
LENGTH: 6372
TYPE: DNA
ORGANISM: Homo Sapiens
US-11-266-748A-30503

Query Match
Best Local Similarity 36.4%; Score 145.6; DB 8; Length 6372;
Matches 202; Conservative 1; Mismatches 70; Indels 5; Gaps 1;

QY 81 ATGCTGTAAATCCAGCACTTCGGAGGCCAAGGTGGGCGGATCACTGAGGTCAAGAGA 140
DB 2899 ACGCTGTAAATCCAGCACTTCGGAGGCCAAGGTGGGCGGATCACTGAGGTCAAGAGA 2840
QY 141 TCGAGACCATCTCGGCCCAACATGTTGAAACCCCGTCTTAAATAATCAAAATATAGC 200
DB 2839 TCAAGACCAAGCTGGCCCAACATGTTGAAACCCCGTCTTAAATAATATTAATTTAGT 2780
QY 201 TGGGCATGTGGGCACACACCTGTAGTCCAGCTACTCAGAGCCGGA-----GATTGAG 255
DB 2779 TGGGGGTGTGGCATGTCTCTGTATGTCAGCTACTAGGGGCGCTGAGGCGAGTTGCTT 2720
QY 256 TGAAGTGAATCGCAGATGAGCCGAAATCAAGATCAAGAGTGAGAGAGTGAGAGC 315
DB 2719 TGACCTGGAGGAGCAGAGGTGGGCCACTTCACTCAGCCTAGGCAAGAGTAAGACTC 2660
QY 316 CGTCTCAAAAACACACAAAAAACAACAAAAACCTAA 333
DB 2659 CAGCTCAAAAAAAGTAAAGCAACAA 2622

RESULT 15
US-11-266-748A-24156/c
Sequence 24156, Application US/11266748A
Publication No. US2006013463A1
GENERAL INFORMATION:
APPLICANT: Harkin, Paul
APPLICANT: Johnston, Patrick
TITLE OF INVENTION: Transcriptome Microarray Technology and
FILE REFERENCE: 55815-0102 (319189)
CURRENT APPLICATION NUMBER: US/11/266,748A
CURRENT FILING DATE: 2005-11-03
PRIOR APPLICATION NUMBER: EP 04105479.2
PRIOR FILING DATE: 2004-11-03
PRIOR APPLICATION NUMBER: EP 04105482.6
PRIOR FILING DATE: 2004-11-03
PRIOR APPLICATION NUMBER: EP 04105483.4
PRIOR FILING DATE: 2004-11-03
PRIOR APPLICATION NUMBER: EP 04105507.0
PRIOR FILING DATE: 2004-11-03
PRIOR APPLICATION NUMBER: EP 04105485.9
PRIOR FILING DATE: 2004-11-03
PRIOR APPLICATION NUMBER: EP 04105484.2
PRIOR FILING DATE: 2004-11-03
PRIOR APPLICATION NUMBER: US 60/662,276
PRIOR FILING DATE: 2005-03-14
PRIOR APPLICATION NUMBER: US 60/700,293
PRIOR FILING DATE: 2005-07-18
NUMBER OF SEQ ID NOS: 483996
SOFTWARE: PatentIn version 3.3
SEQ ID NO 24156
LENGTH: 3324
TYPE: DNA
ORGANISM: Homo Sapiens
US-11-266-748A-24156
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Query Match
Best Local Similarity 36.3%; Score 145; DB 8; Length 3324;
Matches 172; Conservative 0; Mismatches 45; Indels 0; Gaps 0;

QY 81 ATGCTGTAAATCCAGCACTTCGGAGGCCAAGGTGGGCGGATCACTGAGGTCAAGAGA 140
DB 389 ATGCTGTAAATCCAGCACTTCGGAGGCCAAGGTGGGCGGATCACTGAGGTCAAGAGA 330
QY 141 TCGAGACCATCTCGGCCCAACATGTTGAAACCCCGTCTTAAATAATCAAAATATAGC 200
DB 329 TCAAGACCAAGCTGGGAAACATGTTGAAACCCCGTCTTAAATAATCAAAATATAGC 270
QY 201 TGGGCATGTGGGCACACACCTGTAGTCCAGCTACTCAGAGCCGGAAGATTGAGTAGC 260
DB 269 TGGGGGTGTGGCATGTCTCTGTATGTCAGCTACTAGGGGCGCTGAGGCGAGTTGCTT 210
QY 261 TGAAGTGAATCGCAGATGAGCCGAAATCAAGATCAAGATCAAGAGA 297
DB 209 CTTGAACCAAGAGGTGGAGGCTGAGTGAAGCCGAGA 173

RESULT 16
US-11-266-748A-390100
Sequence 390100, Application US/11266748A
Publication No. US2006013463A1
GENERAL INFORMATION:
APPLICANT: Harkin, Paul
APPLICANT: Johnston, Patrick
TITLE OF INVENTION: Transcriptome Microarray Technology and
FILE REFERENCE: 55815-0102 (319189)
CURRENT APPLICATION NUMBER: US/11/266,748A
CURRENT FILING DATE: 2005-11-03
PRIOR APPLICATION NUMBER: EP 04105479.2
PRIOR FILING DATE: 2004-11-03
PRIOR APPLICATION NUMBER: EP 04105482.6
PRIOR FILING DATE: 2004-11-03
PRIOR APPLICATION NUMBER: EP 04105483.4
PRIOR FILING DATE: 2004-11-03
PRIOR APPLICATION NUMBER: EP 04105507.0
PRIOR FILING DATE: 2004-11-03
PRIOR APPLICATION NUMBER: EP 04105485.9
PRIOR FILING DATE: 2004-11-03
PRIOR APPLICATION NUMBER: EP 04105484.2
PRIOR FILING DATE: 2004-11-03
PRIOR APPLICATION NUMBER: US 60/662,276
PRIOR FILING DATE: 2005-03-14
PRIOR APPLICATION NUMBER: US 60/700,293
PRIOR FILING DATE: 2005-07-18
NUMBER OF SEQ ID NOS: 483996
SOFTWARE: PatentIn version 3.3
SEQ ID NO 390100
LENGTH: 1000
TYPE: DNA
ORGANISM: Homo Sapiens
US-11-266-748A-390100

Query Match
Best Local Similarity 36.2%; Score 144.8; DB 8; Length 1000;
Matches 218; Conservative 1; Mismatches 93; Indels 8; Gaps 1;

QY 33 ACCCAATATTAATTAAGACATGTCAGGCCAGGATGACACTGCTGAATGCTGTATC 92
DB 434 ACACATTTTACATTAAGAAATACATGTCAGGCCATGCTGTGCTCAGGCTGTATTC 493
QY 93 CCAAGACTTGGGAGGCCAAGGTGGGCGGATCAGTCAAGATGAGATGAGACCATCC 152
DB 494 CCAGCATTTGGGAGGCCGAGGTGGGCGATCACCGAGGTGAGAGTTCAAGACAGCC 553
QY 153 TGGCCAAATGATGTAACCCCGTCTTAAATAATCAAAATATAGCTGGGCGATGTGG 212
```



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APPLICANT: Mulligan, Karl
TITLE OF INVENTION: Transcription Microarray Technology and
FILE REFERENCE: 55815-0102 (319189)
CURRENT FILING DATE: 2005-11-03
PRIOR APPLICATION NUMBER: US/11/266,748A
PRIOR FILING DATE: 2004-11-03
PRIOR APPLICATION NUMBER: EP 04105479.2
PRIOR FILING DATE: 2004-11-03
PRIOR APPLICATION NUMBER: EP 04105482.6
PRIOR FILING DATE: 2004-11-03
PRIOR APPLICATION NUMBER: EP 04105483.4
PRIOR FILING DATE: 2004-11-03
PRIOR APPLICATION NUMBER: EP 04105507.0
PRIOR FILING DATE: 2004-11-03
PRIOR APPLICATION NUMBER: EP 04105485.9
PRIOR FILING DATE: 2004-11-03
PRIOR APPLICATION NUMBER: EP 04105484.2
PRIOR FILING DATE: 2004-11-03
PRIOR APPLICATION NUMBER: US 60/662,276
PRIOR FILING DATE: 2005-03-14
PRIOR APPLICATION NUMBER: US 60/700,293
PRIOR FILING DATE: 2005-07-18
NUMBER OF SEQ ID NOS: 483996
SOFTWARE: PatentIn version 3.3
SEQ ID NO 206136
LENGTH: 1000
TYPE: DNA
ORGANISM: Homo Sapiens
US-11-266-748A-206136

Query Match      36.1%; Score 144.2; DB 8; Length 1000;
Best Local Similarity 70.0%; Pred. No. 1.9e-21;
Matches 194; Conservative 0; Mismatches 83; Indels 0; Gaps 0;

QY 81 ATGCTGTAAATCCGACACTTCGGAGGCGCAAGGTGGCGGATCACTGAGGTCAAGGA 140
DB 850 ACGCTTAATCCGACACTTCGGAGGCGCAAGGTGGCGGATCACTGAGGTCAAGGA 791
QY 141 TCGGACCATCTGCGCAACATGTGGAACCCCGCTTACTAAATAATCAAAAATATGC 200
DB 790 TCCAGACCAAGCTTCGCAACATGTGGAACCCCGCTTACTAAATAATCAAAAATATGC 731
QY 201 TGGGATGTGGGACACACCTGTAGTCCAGCTACTCAGAGCCGGAATTTGAGTGC 260
DB 730 TGGGATGTGGGACACATGCTGTATCTCACTATTGGGAGGCGTGAAGAGAAATG 671
QY 261 TGAATCCAGAGTGAGCCGGAATATCAAGATCAAGAGTGAGAGACKCCGCT 320
DB 670 CTGGAACCCAGAGGTGAAGGTGAGTGAACCAAGATTGGCGCTGCACTCCAGCATGG 611
QY 321 CAAAACAACAACAACAAAACAAAACCAATAGACA 357
DB 610 GCAAGACAGAAACTCTGTCAAAAAAATAAAAAA 574

RESULT 20
US-11-266-748A-60135/C
Sequence 60135, Application US/11266748A
Publication No. US2006013463A1
GENERAL INFORMATION:
APPLICANT: Harkin, Paul
APPLICANT: Johnston, Patrick
APPLICANT: Mulligan, Karl
TITLE OF INVENTION: Transcription Microarray Technology and
FILE REFERENCE: 55815-0102 (319189)
CURRENT FILING DATE: US/11/266,748A
PRIOR APPLICATION NUMBER: US/11-03
PRIOR FILING DATE: 2005-11-03
PRIOR APPLICATION NUMBER: EP 04105479.2
PRIOR FILING DATE: 2004-11-03
PRIOR APPLICATION NUMBER: EP 04105482.6
PRIOR FILING DATE: 2004-11-03
PRIOR APPLICATION NUMBER: EP 04105483.4
```

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PRIOR FILING DATE: 2004-11-03
PRIOR APPLICATION NUMBER: EP 04105507.0
PRIOR FILING DATE: 2004-11-03
PRIOR APPLICATION NUMBER: EP 04105485.9
PRIOR FILING DATE: 2004-11-03
PRIOR APPLICATION NUMBER: EP 04105484.2
PRIOR FILING DATE: 2004-11-03
PRIOR APPLICATION NUMBER: US 60/662,276
PRIOR FILING DATE: 2005-03-14
PRIOR APPLICATION NUMBER: US 60/700,293
PRIOR FILING DATE: 2005-07-18
NUMBER OF SEQ ID NOS: 483996
SOFTWARE: PatentIn version 3.3
SEQ ID NO 60135
LENGTH: 227968
TYPE: DNA
ORGANISM: Homo Sapiens
US-11-266-748A-60135

Query Match      36.1%; Score 144.2; DB 8; Length 227968;
Best Local Similarity 73.6%; Pred. No. 2.5e-21;
Matches 217; Conservative 1; Mismatches 59; Indels 18; Gaps 2;

QY 81 ATGCTGTAAATCCGACACTTCGGAGGCGCAAGGTGGCGGATCACTGAGGTCAAGGA 140
DB 204862 ACGCTTAATCCGACACTTCGGAGGCGCAAGGTGGCGGATCACTGAGGTCAAGGA 204803
QY 141 TCGGACCATCTGCGCAACATGTGGAACCCCGCTTACTAAATAATCAAAAATATGC 200
DB 204802 TCGGACCATCTGCGCAACATGTGGAACCCCGCTTACTAAATAATCAAAAATATGC 204743
QY 201 TGGGATGTGGGACACACCTGTAGTCCAGCTACTCAGGA-----GCCGGAATTC 253
DB 204742 CGGCGTGTGGGACACATGCTGTATCCAGCTACTCAGGAAGTGAAGGAGAAATGC 204683
QY 254 AGTAGCTAGATTCAGAGTGAGCCGAATCACAATCA-----CAAGTGAG 302
DB 204682 CTGGAACCGGGGAAGCGAGGTTGCAGTGAGCCAAAGATCAACCACTGCATCTTAGGTAA 204623
QY 303 CAGAGTGAGACKCGCTCAAAAACAACAACAAAACCAATAGACA 357
DB 204622 CAGAGTGAGCTCTGTCTCAAAAAAATAAAAAAATAAAAAA 204568

RESULT 21
US-11-266-748A-198112/C
Sequence 198112, Application US/11266748A
Publication No. US2006013463A1
GENERAL INFORMATION:
APPLICANT: Harkin, Paul
APPLICANT: Johnston, Patrick
APPLICANT: Mulligan, Karl
TITLE OF INVENTION: Transcription Microarray Technology and
FILE REFERENCE: 55815-0102 (319189)
CURRENT FILING DATE: US/11/266,748A
PRIOR APPLICATION NUMBER: US/11-03
PRIOR FILING DATE: 2005-11-03
PRIOR APPLICATION NUMBER: EP 04105479.2
PRIOR FILING DATE: 2004-11-03
PRIOR APPLICATION NUMBER: EP 04105482.6
PRIOR FILING DATE: 2004-11-03
PRIOR APPLICATION NUMBER: EP 04105483.4
PRIOR FILING DATE: 2004-11-03
PRIOR APPLICATION NUMBER: EP 04105507.0
PRIOR FILING DATE: 2004-11-03
PRIOR APPLICATION NUMBER: EP 04105485.9
PRIOR FILING DATE: 2004-11-03
PRIOR APPLICATION NUMBER: US 60/662,276
PRIOR FILING DATE: 2005-03-14
PRIOR APPLICATION NUMBER: US 60/700,293
PRIOR FILING DATE: 2005-07-18
```

NUMBER OF SEQ ID NOS: 483996
SOFTWARE: Patentin version 3.3
SEQ ID NO 198112
LENGTH: 1000
TYPE: DNA
ORGANISM: Homo Sapiens
US-11-266-748A-198112

Query Match 35.8%; Score 143; DB 8; Length 1000;
Best Local Similarity 77.6%; Pred. No. 3.3e-21;
Matches 173; Conservative 0; Mismatches 50; Indels 0; Gaps 0;

QY 81 ATGCTGTATCCAGCACTTCGGAGGCCAAGGTGGGGATCACTGAGTCAAGAGA 140
DB 725 ATGCTGTATCCAGCACTTTAGAGGCCGAGGTGGGGATCACTGAGTCAAGAG 666
QY 141 TCGAGACCATCTGGCCAAATGTTGAAACCCGCTTTACTTAAATAAATAATAC 200
DB 665 TCGAGACCATCTGGCCAAATGTTGAAACCCGCTTTACTTAAATAAATAATAC 606
QY 201 TGGGATGTGGCAGACACCTGTAGTCCAGTACTCAGAGCCGAGATTGCACTGAGC 260
DB 605 TGGGATGTGGCAGACACCTGTAGTCTCAGTACTCAGAGCCGAGATTGCACTGAG 546
QY 261 TGAGATCGCAGAGTGGCCGAATCAAGATCAAGATCAAGATGAGC 303
DB 545 CTGTAACCGAGACGAGGTTGCACTGAGCCAGATCTGTC 503

RESULT 22

US-11-266-748A-59101/c
Sequence 59101, Application US/11266748A
Publication No. US20060134663A1

GENERAL INFORMATION:
APPLICANT: Harkin, Paul
APPLICANT: Johnston, Patrick
APPLICANT: Mulligan, Karl
TITLE OF INVENTION: Transcriptome Microarray Technology and
FILE REFERENCE: 55815-0102 (319189)
CURRENT FILING DATE: 2005-11-03
PRIOR FILING DATE: 2004-11-03
PRIOR APPLICATION NUMBER: EP 04105479.2
PRIOR FILING DATE: 2004-11-03
PRIOR APPLICATION NUMBER: EP 04105482.6
PRIOR FILING DATE: 2004-11-03
PRIOR APPLICATION NUMBER: EP 04105483.4
PRIOR FILING DATE: 2004-11-03
PRIOR APPLICATION NUMBER: EP 04105507.0
PRIOR FILING DATE: 2004-11-03
PRIOR APPLICATION NUMBER: EP 04105485.9
PRIOR FILING DATE: 2004-11-03
PRIOR APPLICATION NUMBER: EP 04105484.2
PRIOR FILING DATE: 2004-11-03
PRIOR APPLICATION NUMBER: US 60/662,276
PRIOR FILING DATE: 2005-03-14
PRIOR APPLICATION NUMBER: US 60/700,293
PRIOR FILING DATE: 2005-07-18
NUMBER OF SEQ ID NOS: 483996
SOFTWARE: Patentin version 3.3
SEQ ID NO 59101
LENGTH: 96217
TYPE: DNA
ORGANISM: Homo Sapiens
US-11-266-748A-59101

Query Match 35.8%; Score 143; DB 8; Length 96217;
Best Local Similarity 88.6%; Pred. No. 4.2e-21;
Matches 155; Conservative 0; Mismatches 20; Indels 0; Gaps 0;

QY 81 ATGCTGTATCCAGCACTTCGGAGGCCAAGGTGGGGATCACTGAGTCAAGAGA 140
DB 25922 ACGCTGTATCCAGCACTTTGGAGGCCGAGGTGGGTGATCACTGAGTCAAGAGT 25863

QY 141 TCGAGACCATCTGGCCAAATGTTGAAACCCGCTTTACTTAAATAAATAATAC 200
DB 25862 TCGAGACCATCTGGCCAAATGTTGAAACCCGCTTTACTTAAATAAATAATAC 25803
QY 201 TGGGATGTGGCAGACACCTGTAGTCCAGTACTCAGAGCCGAGATTGCACTGAG 255
DB 25802 TGGGATGTGGTGAACCGCTGTATCCAGTACTCAGAGACCTGAGCAGAG 25748

RESULT 23

US-11-266-748A-27021
Sequence 27021, Application US/11266748A
Publication No. US20060134663A1

GENERAL INFORMATION:
APPLICANT: Harkin, Paul
APPLICANT: Johnston, Patrick
APPLICANT: Mulligan, Karl
TITLE OF INVENTION: Transcriptome Microarray Technology and
FILE REFERENCE: 55815-0102 (319189)
CURRENT FILING DATE: 2005-11-03
PRIOR FILING DATE: 2004-11-03
PRIOR APPLICATION NUMBER: EP 04105479.2
PRIOR FILING DATE: 2004-11-03
PRIOR APPLICATION NUMBER: EP 04105482.6
PRIOR FILING DATE: 2004-11-03
PRIOR APPLICATION NUMBER: EP 04105483.4
PRIOR FILING DATE: 2004-11-03
PRIOR APPLICATION NUMBER: EP 04105507.0
PRIOR FILING DATE: 2004-11-03
PRIOR APPLICATION NUMBER: EP 04105485.9
PRIOR FILING DATE: 2004-11-03
PRIOR APPLICATION NUMBER: EP 04105484.2
PRIOR FILING DATE: 2004-11-03
PRIOR APPLICATION NUMBER: US 60/662,276
PRIOR FILING DATE: 2005-03-14
PRIOR APPLICATION NUMBER: US 60/700,293
PRIOR FILING DATE: 2005-07-18
NUMBER OF SEQ ID NOS: 483996
SOFTWARE: Patentin version 3.3
SEQ ID NO 27021
LENGTH: 1945
TYPE: DNA
ORGANISM: Homo Sapiens
US-11-266-748A-27021

Query Match 35.7%; Score 142.6; DB 8; Length 1945;
Best Local Similarity 69.4%; Pred. No. 4.2e-21;
Matches 209; Conservative 1; Mismatches 85; Indels 6; Gaps 1;

QY 63 AGGATGACACTGCTGATATCCAGCACTTCGGAGGCCAAGGTGGGCGGA 122
DB 1636 AGGCGGGCGTGTGCTATCTGTATCCAGCACTTTGGGTGAGTGGAGCG 1695
QY 123 TCACCTGAGTCAAGAGATCGAACATCTGGCCAAATGTAACCCGCTTTACT 182
DB 1696 TCACCTGAGTCAAGAGATCGAACATCTGGCTTAACTGTGTAACCCGCTTACT 1755
QY 183 AAAAATCAAAAATAGCTGGGCAATGTCAGACACCTGTAGTCCAGTACTCAGAG 242
DB 1756 AAAAATCAAAAATAGCTGGGCAATGTCAGACACCTGTAGTCCAGTACTCAGAG 1815
QY 243 CCGAGATTGCAAGTGAATGATC-----GCAGAGTGAACCGAAATCAAGATCAAG 296
DB 1816 GCTGAGGAGGAATGAGCGTGAACCCGAGAGTGAAGTGGCGCATCGACCTCAGGCT 1875
QY 297 AGTGAAGAGTGAAGCCGCTTCAAAAACAAACAAACAAACAAACAAACAAAC 356
DB 1876 GGGCAACAGACGAAGTCCGTCAAAAACAAACAAACAAACAAACAAACAAAC 1935
QY 357 A 357

Db 1936 A 1936

RESULT 24
US-11-266-748A-58369
Sequence 58369, Application US/11266748A
Publication No. US2006013463A1
GENERAL INFORMATION:
APPLICANT: Harkin, Paul
APPLICANT: Johnston, Patrick
APPLICANT: Mulligan, Karl
TITLE OF INVENTION: Transcriptome Microarray Technology and
FILE REFERENCE: 55815-0102 (319189)
CURRENT FILING DATE: 2005-11-03
PRIOR FILING DATE: 2005-11-03
PRIOR APPLICATION NUMBER: EP 04105479.2
PRIOR FILING DATE: 2004-11-03
PRIOR APPLICATION NUMBER: EP 04105482.6
PRIOR FILING DATE: 2004-11-03
PRIOR APPLICATION NUMBER: EP 04105483.4
PRIOR FILING DATE: 2004-11-03
PRIOR APPLICATION NUMBER: EP 04105507.0
PRIOR FILING DATE: 2004-11-03
PRIOR APPLICATION NUMBER: EP 04105485.9
PRIOR FILING DATE: 2004-11-03
PRIOR APPLICATION NUMBER: EP 04105484.2
PRIOR FILING DATE: 2004-11-03
PRIOR APPLICATION NUMBER: US 60/662,276
PRIOR FILING DATE: 2005-03-14
PRIOR APPLICATION NUMBER: US 60/700,293
PRIOR FILING DATE: 2005-07-18
NUMBER OF SEQ ID NOS: 483996
SOFTWARE: PatentIn version 3.3
SEQ ID NO 58369
LENGTH: 1945
TYPE: DNA
ORGANISM: Homo Sapiens
US-11-266-748A-58369

Query Match 35.7%; Score 142.6; DB 8; Length 1945;
Best Local Similarity 69.4%; Pred. No. 4.2e-21;
Matches 209; Conservative 1; Mismatches 85; Indels 6; Gaps 1;

63 AGGATGACACTGCTGATGCTTTATCCAGACATTCGGGAGGCCAAGTGGCGGA 122
1636 AGGCGGCGCTGTGCTGCTGATGCCAGACATTTGGTGCTGAGGTGACGGA 1695
123 TCACCTGAGTCAAGAGATCGAGCCATCTGGCCAAATGGTGAACCCCGCTTTACT 182
1696 TCACCTGAGTCAAGAGATTTGAGACCATCTGGCTAAGCTGTGAACCCCGCTTACT 1755
183 AAAAATACAAAAATAGCTGGGCGATGTGGCACACACCTGTGATCCAGCTACTCAGAG 242
1756 AAAAATACAAAAATAGCGGGCGCTGGGGGTGGCGCGCTGTGATCCAGCTACTCAGAG 1815
243 CCGGAGATTGCACTGAGCTGAGATC-----GCAAGTGAAGCCGAAATCAGATCAAG 296
1816 GCTGGGCGAGGAGAAATGCGTGAACCCCGGAGTCAAGATCGGCACTGCACTCAGCT 1875
297 AGTGAGCAGAGTGAACKCCGCTCTCAAAAAACAACAACAAAAAACAATAGAGC 356
1876 GGGCAACAGAGCAAGAGTCCGCTCTCAAAAAAAGAAAAAAGAAAAAAGAAAAA 1935
357 A 357
1936 A 1936

RESULT 25
US-11-266-748A-28208
Sequence 28208, Application US/11266748A
Publication No. US2006013463A1

GENERAL INFORMATION:
APPLICANT: Harkin, Paul
APPLICANT: Johnston, Patrick
APPLICANT: Mulligan, Karl
TITLE OF INVENTION: Transcriptome Microarray Technology and
FILE REFERENCE: 55815-0102 (319189)
CURRENT FILING DATE: 2005-11-03
PRIOR FILING DATE: 2005-11-03
PRIOR APPLICATION NUMBER: EP 04105479.2
PRIOR FILING DATE: 2004-11-03
PRIOR APPLICATION NUMBER: EP 04105482.6
PRIOR FILING DATE: 2004-11-03
PRIOR APPLICATION NUMBER: EP 04105483.4
PRIOR FILING DATE: 2004-11-03
PRIOR APPLICATION NUMBER: EP 04105507.0
PRIOR FILING DATE: 2004-11-03
PRIOR APPLICATION NUMBER: EP 04105485.9
PRIOR FILING DATE: 2004-11-03
PRIOR APPLICATION NUMBER: EP 04105484.2
PRIOR FILING DATE: 2004-11-03
PRIOR APPLICATION NUMBER: US 60/662,276
PRIOR FILING DATE: 2005-03-14
PRIOR APPLICATION NUMBER: US 60/700,293
PRIOR FILING DATE: 2005-07-18
NUMBER OF SEQ ID NOS: 483996
SOFTWARE: PatentIn version 3.3
SEQ ID NO 28208
LENGTH: 1421559
TYPE: DNA
ORGANISM: Homo Sapiens
FEATURE:
NAME/KEY: misc feature
LOCATION: (41394)..(41394)
OTHER INFORMATION: n is a, c, g, or t
FEATURE:
NAME/KEY: misc feature
LOCATION: (83609)..(83609)
OTHER INFORMATION: n is a, c, g, or t
FEATURE:
NAME/KEY: misc feature
LOCATION: (1022504)..(1022504)
OTHER INFORMATION: n is a, c, g, or t
FEATURE:
NAME/KEY: misc feature
LOCATION: (1041105)..(1041105)
OTHER INFORMATION: n is a, c, g, or t
FEATURE:
NAME/KEY: misc feature
LOCATION: (1122667)..(1122667)
OTHER INFORMATION: n is a, c, g, or t
US-11-266-748A-28208

Query Match 35.7%; Score 142.6; DB 8; Length 1421559;
Best Local Similarity 73.0%; Pred. No. 5.8e-22;
Matches 195; Conservative 1; Mismatches 70; Indels 1; Gaps 1;

85 CTGTAATCCGACACTTCGGGAGGCGAAGGTGGGCGATCACTGAGGTCAAGATGCA 144
1069903 CTGTAATCCGACACTTCGGGAGGCGAAGGTGGGCGATCACTGAGGTCAAGATGCA 1069962
145 GACCATCTGGCCCAACATGTGTAACCCGCTTTACTAAATAAATAAATAATAGCTGG 204
1069963 GACCATCTGGCCCAACATGTGTAACCCGCTTTACTAAATAAATAAATAATAGCTGG 1070022
205 CATGTGGCACACACTGTAGTCCAGCTACTCAGGA-GCCGAGATTGCACTGAGCTGA 263
1070023 TTTGGTGTGTGTGCTGTAGTCCAGCTACTCAGGAGGCTGAAGCAAGAGATGCTTG 1070082
264 GATCGAGAGTGAAGCCCAATATCAGATCAAGAGTGAAGAGAGAGAGAGAGAGAGAG 323
1070083 AATCGGAGGCGGAGGTGCAAGTCTTGAAGATGAGTGAAGAGAGAGAGAGAGAG 1070142


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; PRIOR APPLICATION NUMBER: EP 04105483.4
; PRIOR FILING DATE: 2004-11-03
; PRIOR APPLICATION NUMBER: EP 04105507.0
; PRIOR FILING DATE: 2004-11-03
; PRIOR APPLICATION NUMBER: EP 04105485.9
; PRIOR FILING DATE: 2004-11-03
; PRIOR APPLICATION NUMBER: EP 04105484.2
; PRIOR FILING DATE: 2004-11-03
; PRIOR APPLICATION NUMBER: US 60/662,276
; PRIOR FILING DATE: 2005-03-14
; PRIOR APPLICATION NUMBER: US 60/700,293
; PRIOR FILING DATE: 2005-07-18
; NUMBER OF SEQ ID NOS: 483996
; SOFTWARE: PatentIn version 3.3
; SEQ ID NO: 293957
; LENGTH: 1000
; TYPE: DNA
; ORGANISM: Homo Sapiens
US-11-266-748A-293957

Query Match      35.6%; Score 142.4; DB 8; Length 1000;
Best Local Similarity 73.0%; Pred. No. 4.5e-21;
Matches 219; Conservative 1; Mismatches 57; Indels 23; Gaps 2;

OY 81 ATGCTGTAATCCGACACTTCGGAGGCGCAAGTGGGGGATCCAGTCAAGTCAAGAGA 140
DB 133 ATGCTGTAATCCGACACTTCGGAGGCGCAAGTGGGGGATCCAGTCAAGTCAAGAGA 192
OY 141 TCGAGACCATCTGCGCCAAATGTAAGAAACCCGCTTTTACTTAAATAACAAAAATAGC 200
DB 193 TCGAGACCATCTGCGCCAAATGTAAGAAACCCGCTTTTACTTAAATAACAAAAATAGC 252
OY 201 TGGGATGATGCGACACACCTGTACTCCAGTACTTCAAGAGAGAGAGAGAGAGAGAG 253
DB 253 TGGGATGATGCGACACACCTGTACTCCAGTACTTCAAGAGAGAGAGAGAGAGAGAG 312
OY 254 AGTGAAGTGAATCCAGAGTGAAGGCGCAATCAGAGATCA-----CAGA 297
DB 313 CTGTAAGTGGGGAGGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAG 372
OY 298 GTGAGCAGAGTGAACCKCGCTCTCAAAAACAAACAAACAAACAAACAAACCAATAGACA 357
DB 373 GGTGACAGAGTGAATCTCATCTCAAAAAACAAACAAACAAACAAACCAATAGACA 432

RESULT 29
US-11-266-748A-343660
; Sequence 343660, Application US/11266748A
; Publication No. US2006013463A1
; GENERAL INFORMATION:
; APPLICANT: Harkin, Paul
; APPLICANT: Johnston, Patrick
; TITLE OF INVENTION: Transcriptome Microarray Technology and
; FILE REFERENCE: 55815-0102 (319189)
; CURRENT APPLICATION NUMBER: US/11/266,748A
; PRIOR FILING DATE: 2005-11-03
; PRIOR APPLICATION NUMBER: EP 04105479.2
; PRIOR FILING DATE: 2004-11-03
; PRIOR APPLICATION NUMBER: EP 04105482.6
; PRIOR FILING DATE: 2004-11-03
; PRIOR APPLICATION NUMBER: EP 04105483.4
; PRIOR FILING DATE: 2004-11-03
; PRIOR APPLICATION NUMBER: EP 04105507.0
; PRIOR FILING DATE: 2004-11-03
; PRIOR APPLICATION NUMBER: EP 04105485.9
; PRIOR FILING DATE: 2004-11-03
; PRIOR APPLICATION NUMBER: EP 04105484.2
; PRIOR FILING DATE: 2004-11-03
; PRIOR APPLICATION NUMBER: US 60/662,276
; PRIOR FILING DATE: 2005-03-14
; PRIOR APPLICATION NUMBER: US 60/700,293
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; PRIOR FILING DATE: 2005-07-18
; NUMBER OF SEQ ID NOS: 483996
; SOFTWARE: PatentIn version 3.3
; SEQ ID NO 343660
; LENGTH: 1000
; TYPE: DNA
; ORGANISM: Homo Sapiens
US-11-266-748A-343660

Query Match      35.6%; Score 142.4; DB 8; Length 1000;
Best Local Similarity 69.9%; Pred. No. 4.5e-21;
Matches 207; Conservative 1; Mismatches 82; Indels 6; Gaps 1;

OY 85 CTGTAATCCGACACTTCGGAGGCGCAAGTGGGGGATCCAGTCAAGTCAAGAGA 144
DB 535 CTGTAATCCGACACTTCGGAGGCGCAAGTGGGGGATCCAGTCAAGTCAAGAGA 594
OY 145 GACCATCTGCGCCAAATGTAAGAAACCCGCTTTTACTTAAATAACAAAAATAGCTGGG 204
DB 595 GACCATCTGCGCCAAATGTAAGAAACCCGCTTTTACTTAAATAACAAAAATAGCTGGG 654
OY 205 CATGTGCGACACACTGTACTCCAGTACTTCAAGAGCGCGAGATTCAGTGAAGTGAAG 264
DB 655 CGTGTGCGACGCGCTGTACTCCAGTACTTCAAGAGCGCGAGATTCAGTGAAGTGAAG 714
OY 265 ATCGAGAGTGAAGCGCAATCAGAGATCA-----CAGAGTGAAGCAGAGTGAAGCCKCGT 318
DB 715 AACCGAGAGCGAGATCATCTGCACTCCAGCGCTGGGTGACAGAGTGAAGTGAAGTCTGT 774
OY 319 CTCAAAAACAAACAAACAAACAAACAAACCAATAGACATTCGATTCGCGGTTTC 374
DB 775 CTCAAAAACAAACAAACAAACAAACAAACCAATAGACATTCGATTCGCGGTTTC 830

RESULT 30
US-11-266-748A-345386/c
; Sequence 345386, Application US/11266748A
; Publication No. US2006013463A1
; GENERAL INFORMATION:
; APPLICANT: Harkin, Paul
; APPLICANT: Johnston, Patrick
; TITLE OF INVENTION: Transcriptome Microarray Technology and
; FILE REFERENCE: 55815-0102 (319189)
; CURRENT APPLICATION NUMBER: US/11/266,748A
; PRIOR FILING DATE: 2005-11-03
; PRIOR APPLICATION NUMBER: EP 04105479.2
; PRIOR FILING DATE: 2004-11-03
; PRIOR APPLICATION NUMBER: EP 04105482.6
; PRIOR FILING DATE: 2004-11-03
; PRIOR APPLICATION NUMBER: EP 04105483.4
; PRIOR FILING DATE: 2004-11-03
; PRIOR APPLICATION NUMBER: EP 04105507.0
; PRIOR FILING DATE: 2004-11-03
; PRIOR APPLICATION NUMBER: EP 04105485.9
; PRIOR FILING DATE: 2004-11-03
; PRIOR APPLICATION NUMBER: EP 04105484.2
; PRIOR FILING DATE: 2004-11-03
; PRIOR APPLICATION NUMBER: US 60/662,276
; PRIOR FILING DATE: 2005-03-14
; PRIOR APPLICATION NUMBER: US 60/700,293
; PRIOR FILING DATE: 2005-07-18
; NUMBER OF SEQ ID NOS: 483996
; SOFTWARE: PatentIn version 3.3
; SEQ ID NO 345386
; LENGTH: 1000
; TYPE: DNA
; ORGANISM: Homo Sapiens
US-11-266-748A-345386

Query Match      35.6%; Score 142.4; DB 8; Length 1000;
Best Local Similarity 73.0%; Pred. No. 4.5e-21;
```


Matches 219; Conservative 1; Mismatches 57; Indels 23; Gaps 2;

QY 81 ATGCCGTAATCCAGCACTTGGGAGGCGCAAGTGGGGGATTCACCTGAGGTCAAGACA 140
DB 868 ATGCTGTATCCAGCACTTAAGGAGGCGGAGGAGGATCTCCAGAGGTCAAGAGT 809
QY 141 TCGAGACCATCTGGCCCAATGATGTAACCCCGTCTTACTTAATAAATAAATAATAC 200
DB 808 TCGAGACCAAGCTGGCCCAATGATGTAACCCCGTCTTACTTAATAAATAAATAATAC 749
QY 201 TGGGCAATGTGGCAACACCTGTAGTCCAGTACTCAGAA-----GCCGAGATTGC 253
DB 748 TGGGCAATGTGGCAACACCTGTAGTCCAGTACTCAGAA-----GCCGAGATTGC 689
QY 254 AGTGAGCTGAGATCCAGAGTGAAGCCGAATCAAGATCA-----CAGA 297
DB 668 CTTGAACCTGGGAGGCGAGGTTGACGTAGACAAGATCACTGACTGACAGCCCTG 629
QY 238 GTGACGAGAGTGAACKCCGCTCAAAAAACAACAACAAAAAACATAAGACA 357
DB 628 GGTGACAGAGTATCTCATCTCAAAAAAAAAAAAAAAAAAAAAAAAAAAAAA 569

RESULT 31
US-11-266-748A-403868/c
Sequence 403868, Application US/11266748A
Publication No. US20060134663A1
GENERAL INFORMATION:
APPLICANT: Harkin, Paul
APPLICANT: Johnston, Patrick
TITLE OF INVENTION: Transcriptome Microarray Technology and
FILE REFERENCE: 55815-0102 (319189)
CURRENT FILING DATE: 2005-11-03
PRIOR APPLICATION NUMBER: US/11/266,748A
PRIOR FILING DATE: 2004-11-03
PRIOR APPLICATION NUMBER: EP 04105479.2
PRIOR FILING DATE: 2004-11-03
PRIOR APPLICATION NUMBER: EP 04105482.6
PRIOR FILING DATE: 2004-11-03
PRIOR APPLICATION NUMBER: EP 04105483.4
PRIOR FILING DATE: 2004-11-03
PRIOR APPLICATION NUMBER: EP 04105507.0
PRIOR FILING DATE: 2004-11-03
PRIOR APPLICATION NUMBER: EP 04105485.9
PRIOR FILING DATE: 2004-11-03
PRIOR APPLICATION NUMBER: EP 04105484.2
PRIOR FILING DATE: 2004-11-03
PRIOR APPLICATION NUMBER: US 60/662,276
PRIOR FILING DATE: 2005-03-14
PRIOR APPLICATION NUMBER: US 60/700,293
PRIOR FILING DATE: 2005-07-18
NUMBER OF SEQ ID NOS: 483996
SOFTWARE: PatentIn version 3.3
SEQ ID NO 403868
LENGTH: 1000
TYPE: DNA
ORGANISM: Homo Sapiens
US-11-266-748A-403868

Query Match 35.6%; Score 142.4; DB 8; Length 1000;
Best Local Similarity 69.9%; Pred. No. 4.5e-21;
Matches 207; Conservative 1; Mismatches 82; Indels 6; Gaps 1;

QY 85 CTGTATATCCAGCACTTGGGAGGCGCAAGTGGGGGATCACTGAGGTCAAGATGA 144
DB 466 CTGTATATCCAGCACTTGGGAGGCGCGAGGCGGATCGATCACTGAGGTCAAGATGA 407
QY 145 GACCATCTGGCCCAATGATGTAACCCCGTCTTACTTAATAAATAAATAATGCTGG 204
DB 406 GACCATCTGGCCCAATGATGTAACCCCGTCTTACTTAATAAATAAATAATGCTGG 347
QY 205 CATGGTGACACACCTGTATGCTCAAGTCAAGGAGCGGAGATTGCAATGAGCTGAG 264

DB 346 CCGTGTGGCAGGCGCTGTATCCAGTACTCGGAGGCTGAGGAGGATGCTTGG 287
QY 245 ATCGAGAGTGAAGCCGAATCAAGATCA-----CAGAGTGAAGAGTGAAGKCCGT 318
DB 246 AACCCAGAGGCGGAGATCATGCTATGCACTGACCTGGGAGTGAAGATGCTGT 227
QY 319 CTCAAAAACAACAACAAAAAACATAAGACATTGTCATCTGCGGCTTC 374
DB 226 CTCAAAAACAACAACAAAAAACATAAGACATTGTCATCTGCGGCTTC 171

RESULT 32
US-11-266-748A-406117
Sequence 406117, Application US/11266748A
Publication No. US20060134663A1
GENERAL INFORMATION:
APPLICANT: Harkin, Paul
APPLICANT: Johnston, Patrick
TITLE OF INVENTION: Transcriptome Microarray Technology and
FILE REFERENCE: 55815-0102 (319189)
CURRENT FILING DATE: 2005-11-03
PRIOR APPLICATION NUMBER: US/11/266,748A
PRIOR FILING DATE: 2004-11-03
PRIOR APPLICATION NUMBER: EP 04105479.2
PRIOR FILING DATE: 2004-11-03
PRIOR APPLICATION NUMBER: EP 04105482.6
PRIOR FILING DATE: 2004-11-03
PRIOR APPLICATION NUMBER: EP 04105483.4
PRIOR FILING DATE: 2004-11-03
PRIOR APPLICATION NUMBER: EP 04105507.0
PRIOR FILING DATE: 2004-11-03
PRIOR APPLICATION NUMBER: EP 04105485.9
PRIOR FILING DATE: 2004-11-03
PRIOR APPLICATION NUMBER: EP 04105484.2
PRIOR FILING DATE: 2004-11-03
PRIOR APPLICATION NUMBER: US 60/662,276
PRIOR FILING DATE: 2005-03-14
PRIOR APPLICATION NUMBER: US 60/700,293
PRIOR FILING DATE: 2005-07-18
NUMBER OF SEQ ID NOS: 483996
SOFTWARE: PatentIn version 3.3
SEQ ID NO 406117
LENGTH: 1000
TYPE: DNA
ORGANISM: Homo Sapiens
US-11-266-748A-406117

Query Match 35.6%; Score 142.4; DB 8; Length 1000;
Best Local Similarity 73.0%; Pred. No. 4.5e-21;
Matches 219; Conservative 1; Mismatches 57; Indels 23; Gaps 2;

QY 81 ATGCCGTAATCCAGCACTTGGGAGGCGCAAGTGGGGGATCACTGAGGTCAAGACA 140
DB 133 ATGCCGTAATCCAGCACTTAAGGAGGCGGAGGAGGATCTCCAGAGGTCAAGAGT 192
QY 141 TCGAGACCATCTGGCCCAATGATGTAACCCCGTCTTACTTAATAAATAAATAATGCTGG 200
DB 193 TCGAGACCAAGCTGGCCCAATGATGTAACCCCGTCTTACTTAATAAATAAATAATGCTGG 252
QY 201 TGGGCAATGTGGCAACACCTGTAGTCCAGTACTCAGAA-----GCCGAGATTGC 253
DB 253 TGGGCAATGTGGCAACACCTGTAGTCCAGTACTCAGAA-----GCCGAGATTGC 312
QY 254 AGTGAGCTGAGATCCAGAGTGAAGCCGAATCAAGATCA-----CAGA 297
DB 313 CTTGAACCTGGGAGGCGAGGTTGACGTAGACAAGATCACTGACTGACAGCCCTG 372
QY 298 GTGACGAGAGTGAACKCCGCTCAAAAAACAACAACAAAAAACATAAGACA 357
DB 373 GGTGACAGAGTATCTCATCTCAAAAAAAAAAAAAAAAAAAAAAAAAAAAAA 432

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RESULT 33
US-11-266-748A-474914
; Sequence 474914, Application US/11266748A
; Publication No. US20060134663A1
; GENERAL INFORMATION:
; APPLICANT: Harkin, Paul
; APPLICANT: Johnston, Patrick
; APPLICANT: Mulligan, Karl
; TITLE OF INVENTION: Transcriptome Microarray Technology and
; FILE REFERENCE: 55815-0102 (319189)
; CURRENT FILING DATE: 2005-11-03
; PRIOR FILING DATE: 2005-11-03
; PRIOR APPLICATION NUMBER: EP 04105482.6
; PRIOR FILING DATE: 2004-11-03
; PRIOR APPLICATION NUMBER: EP 04105483.4
; PRIOR FILING DATE: 2004-11-03
; PRIOR APPLICATION NUMBER: EP 04105507.0
; PRIOR FILING DATE: 2004-11-03
; PRIOR APPLICATION NUMBER: EP 04105485.9
; PRIOR FILING DATE: 2004-11-03
; PRIOR APPLICATION NUMBER: EP 04105484.2
; PRIOR FILING DATE: 2004-11-03
; PRIOR APPLICATION NUMBER: US 60/662,276
; PRIOR FILING DATE: 2005-03-14
; PRIOR APPLICATION NUMBER: US 60/700,293
; PRIOR FILING DATE: 2005-07-18
; NUMBER OF SEQ ID NOS: 483996
; SOFTWARE: PatentIn version 3.3
; SEQ ID NO 474914
; LENGTH: 1000
; TYPE: DNA
; ORGANISM: Homo Sapiens
US-11-266-748A-474914

Query Match      35.6%; Score 142.4; DB 8; Length 1000;
Best Local Similarity 69.9%; Pred. No. 4.5e-21;
Matches 207; Conservative 1; Mismatches 82; Indels 6; Gaps 1;

QY 85 CTGTATCCGACGACCTTCGGAGGCGCAAGGTGGCGGATCACTGAGTCAAGATCGA 144
DB 535 CTGTATCCGACGACCTTCGGAGGCGCAAGGTGGCGGATCACTGAGTCAAGATCGA 594
QY 145 GACCATCTCTGGCCAAATGCTGTAACCCCGCTTTTACTTAAATAACAAAAATAGCTGG 204
DB 595 GACCATCTCTGGCCAAATGCTGTAACCCCGCTTTTACTTAAATAACAAAAATAGCTGG 654
QY 205 CATGTGGGACACACTCTGTAAGTCCCAAGCTACTCAGAGCGCGAGATGTGAGTGA 264
DB 655 CATGTGGGACACACTCTGTAAGTCCCAAGCTACTCAGAGCGCGAGATGTGAGTGA 714
QY 265 ATCGAGAGTGAAGCCGAATTCACAGATCA-----CAGATGAGAGATGAGACCCGT 318
DB 715 AACCCAGAGAGGGAATCTATGCCATTCCTCAGAGCTGGTGAAGATGAAATCTCTGT 774
QY 319 CTCAAAAACAAACAAAAAACAACAAAAACATTAAGACATTTGCTTGGCGTTG 374
DB 775 CTCAAAAAAGAAAAAGAAAAAATTCCTTAAGTGTGCTTGGCCCTC 830

RESULT 34
US-11-266-748A-477163/c
; Sequence 477163, Application US/11266748A
; Publication No. US20060134663A1
; GENERAL INFORMATION:
; APPLICANT: Harkin, Paul
; APPLICANT: Johnston, Patrick
; APPLICANT: Mulligan, Karl
; TITLE OF INVENTION: Transcriptome Microarray Technology and
; FILE REFERENCE: 55815-0102 (319189)
; CURRENT FILING DATE: 2005-11-03
; PRIOR FILING DATE: 2005-11-03
; PRIOR APPLICATION NUMBER: EP 04105482.6
; PRIOR FILING DATE: 2004-11-03
; PRIOR APPLICATION NUMBER: EP 04105483.4
; PRIOR FILING DATE: 2004-11-03
; PRIOR APPLICATION NUMBER: EP 04105507.0
; PRIOR FILING DATE: 2004-11-03
; PRIOR APPLICATION NUMBER: EP 04105485.9
; PRIOR FILING DATE: 2004-11-03
; PRIOR APPLICATION NUMBER: EP 04105484.2
; PRIOR FILING DATE: 2004-11-03
; PRIOR APPLICATION NUMBER: US 60/662,276
; PRIOR FILING DATE: 2005-03-14
; PRIOR APPLICATION NUMBER: US 60/700,293
; PRIOR FILING DATE: 2005-07-18
; NUMBER OF SEQ ID NOS: 483996
; SOFTWARE: PatentIn version 3.3
; SEQ ID NO 477163
; LENGTH: 1000
; TYPE: DNA
; ORGANISM: Homo Sapiens
US-11-266-748A-477163

Query Match      35.6%; Score 142.4; DB 8; Length 1000;
Best Local Similarity 73.0%; Pred. No. 4.5e-21;
Matches 219; Conservative 1; Mismatches 57; Indels 23; Gaps 2;
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FILE REFERENCE: 55815-0102 (319189)
; CURRENT APPLICATION NUMBER: US/11/266,748A
; CURRENT FILING DATE: 2005-11-03
; PRIOR APPLICATION NUMBER: EP 04105479.2
; PRIOR FILING DATE: 2004-11-03
; PRIOR APPLICATION NUMBER: EP 04105482.6
; PRIOR FILING DATE: 2004-11-03
; PRIOR APPLICATION NUMBER: EP 04105483.4
; PRIOR FILING DATE: 2004-11-03
; PRIOR APPLICATION NUMBER: EP 04105507.0
; PRIOR FILING DATE: 2004-11-03
; PRIOR APPLICATION NUMBER: EP 04105485.9
; PRIOR FILING DATE: 2004-11-03
; PRIOR APPLICATION NUMBER: EP 04105484.2
; PRIOR FILING DATE: 2004-11-03
; PRIOR APPLICATION NUMBER: US 60/662,276
; PRIOR FILING DATE: 2005-03-14
; PRIOR APPLICATION NUMBER: US 60/700,293
; PRIOR FILING DATE: 2005-07-18
; NUMBER OF SEQ ID NOS: 483996
; SOFTWARE: PatentIn version 3.3
; SEQ ID NO 477163
; LENGTH: 1000
; TYPE: DNA
; ORGANISM: Homo Sapiens
US-11-266-748A-477163

Query Match      35.6%; Score 142.4; DB 8; Length 1000;
Best Local Similarity 73.0%; Pred. No. 4.5e-21;
Matches 219; Conservative 1; Mismatches 57; Indels 23; Gaps 2;

QY 81 ATGCTGTATCCAGACGACCTTCGGAGGCGCAAGGTGGCGGATCACTGAGTCAAGATCGA 140
DB 868 ATGCTGTATCCAGACGACCTTCGGAGGCGCAAGGTGGCGGATCACTGAGTCAAGATCGA 809
QY 141 TCGAGACCATCTCTGGCCAAATGCTGTAACCCCGCTTTTACTTAAATAACAAAAATAGC 200
DB 808 TCGAGACCATCTCTGGCCAAATGCTGTAACCCCGCTTTTACTTAAATAACAAAAATAGC 749
QY 201 TGGGATGATGGGACACACCTGTAGTCCCAAGCTACTCAGAGTTC 253
DB 748 TGGGATGATGGGACACACCTGTAGTCCCAAGCTACTCAGAGTTC 689
QY 254 AGTGAGTGAATGCGACAGTGAAGCCGAATTCACAGATCA-----CAGA 297
DB 688 CTGGAACCTGGGAGGACAGGTTGAGTGAAGACAGATCAAGTCACTGATCCAGCCTG 629
QY 298 GTGAGCAGAGTGAAGCCKCGCTCTCAAAAACAACAACAAAAAACAATTAAGCA 357
DB 628 GTGAGCAGAGTGAATCTCATCTCAAAAAAAAAAAAAAAAAAAAAA 569

RESULT 35
US-11-191-644-1
; Sequence 1, Application US/1191644
; Publication No. US20060141529A1
; GENERAL INFORMATION:
; APPLICANT: KOLESKE, ANTHONY JOHN
; APPLICANT: BOYLE, SCOTT NILE
; APPLICANT: SCHWEITZER, BARRY
; APPLICANT: MICHAUD, GREG
; APPLICANT: PREDKI, PAUL
; TITLE OF INVENTION: COMPOSITIONS, KITS AND ASSAYS CONTAINING REAGENTS
; FILE REFERENCE: INV-1007-UT
; CURRENT FILING DATE: 2005-07-27
; PRIOR APPLICATION NUMBER: US/11/191,644
; PRIOR FILING DATE: 2004-07-27
; NUMBER OF SEQ ID NOS: 29
; SOFTWARE: PatentIn version 3.3
; SEQ ID NO 1
; LENGTH: 157866
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TYPE: DNA
ORGANISM: Homo sapiens
US-11-191-644-1

Query Match 35.6%; Score 142.2; DB 7; Length 157866;
Best Local Similarity 73.6%; Pred. No. 6.3e-21;
Matches 229; Conservative 1; Mismatches 59; Indels 22; Gaps 3;

QY 81 ATGCTGTATATCCGACACTTGGGAGGCCAAGTGGCGGATCACTGAGTCAAGAGA 140
DB 62586 ATGCTGTATATCCGACACTTGGGAGGCCAAGTGGCGGATCACTGAGTCAAGAGA 62643
QY 141 TCGAGACCATCTGCGCAACATGTAAGAAACCCGCTTACTTAATAAATAATTAATGTC 200
DB 62644 TTGAGACCATCTGCGCAACATGTAAGAAACCCGCTTACTTAATAAATAATTAATGTC 62703
QY 201 TGGGATGATGCGACACACTGTAGTCCAGCTACTCAGCA-----GCCGAGATTGC 253
DB 62704 TGTGATGATGCGACACACTGTAGTCCAGCTACTCAGCA-----GCCGAGATTGC 62763
QY 254 AGTGAAGTGAATGCCAAGTGAAGCCGAATCAAGAT-----CAAGAAGTG 300
DB 62764 CTGTAACCCAGAGGACAGAGTTGCAAGTGAAGTTCACCACTGCACTCCAGACTG 62823
QY 301 AGCAGAGTGAAGCCGCTCTCAAAAAACAACAACAAAAAACAACATTAAGACATTG 360
DB 62824 CTGACAGTGAAGCTCGTCTCAAAAAACAACAACAAAAACAACAACAGAGAGG 62883
QY 361 TCCATCTGCGG 371
DB 62884 ACTGCTTCTG 62894

RESULT 36
US-11-293-697-23/c
Sequence 23, Application US/11293697
Publication No. US20060105376A1

GENERAL INFORMATION:
APPLICANT: HELIX RESEARCH INSTITUTE
TITLE OF INVENTION: Novel full length cDNA
FILE REFERENCE: H1-A0106
CURRENT APPLICATION NUMBER: US/11/293,697
CURRENT FILING DATE: 2005-12-05
PRIOR APPLICATION NUMBER: US/10/108,260
PRIOR FILING DATE: 2002-03-28
NUMBER OF SEQ ID NOS: 5458
SOFTWARE: PatentIn Ver. 2.1
SEQ ID NO 23
LENGTH: 2800
TYPE: DNA
ORGANISM: Homo sapiens
US-11-293-697-23

Query Match 35.5%; Score 142; DB 8; Length 2800;
Best Local Similarity 72.3%; Pred. No. 5.7e-21;
Matches 198; Conservative 1; Mismatches 71; Indels 4; Gaps 1;

QY 76 GCTGAATGCTGTATATCCGACACTTGGGAGGCCAAGTGGCGGATCACTGAGTCA 135
DB 847 GTTTCACGCTGTATATCCGACACTTGGGAGGCCAAGTGGCGGATCACTGAGTCA 788
QY 136 AGAATGAGACCATCTGCGCAACATGTAAGAAACCCGCTTACTTAATAAATAATTAATGTC 195
DB 787 GGAATTCAGACCAAGCCCTGGCCAAATGTGTAACCCGCTTACTTAATAAATAATTAATGTC 728
QY 196 ATACTGGGATGATGAGACACACTGTAGTCCAGCTACTCAGAGCCGAGATTGAG 255
DB 727 TTAAGTGGGCTGTGAGACATGCTGTATATCCAGCTTACCGGAGGCTGAGGAGAG 668
QY 256 TGAAGTGAATGCAAGTGAAGCCGAATCAAGATCAAGAGTGAAGAGAGAGAGC 315
DB 667 AATCACTGAACCCGGAGGCGAGAGGCTC-----CAGTCTAGTCAAAAAAAGCAAGACTC 612

QY 316 CGTCAAAAAACAACAACAAAAAACC 349
DB 611 AGTCTCAAAAAACAACCAAGAGCCAAACAAAAACC 578

RESULT 37
US-11-293-697-604/c
Sequence 604, Application US/11293697
Publication No. US20060105376A1

GENERAL INFORMATION:
APPLICANT: HELIX RESEARCH INSTITUTE
TITLE OF INVENTION: Novel full length cDNA
FILE REFERENCE: H1-A0106
CURRENT APPLICATION NUMBER: US/11/293,697
CURRENT FILING DATE: 2005-12-05
PRIOR APPLICATION NUMBER: US/10/108,260
PRIOR FILING DATE: 2002-03-28
NUMBER OF SEQ ID NOS: 5458
SOFTWARE: PatentIn Ver. 2.1
SEQ ID NO 604
LENGTH: 3252
TYPE: DNA
ORGANISM: Homo sapiens
US-11-293-697-604

Query Match 35.5%; Score 142; DB 8; Length 3252;
Best Local Similarity 77.5%; Pred. No. 5.7e-21;
Matches 172; Conservative 0; Mismatches 50; Indels 0; Gaps 0;

QY 81 ATGCTGTATATCCGACACTTGGGAGGCCAAGTGGCGGATCACTGAGTCAAGAGA 140
DB 2938 AGCTGTATATCCGACACTTGGGAGGCCAAGTGGCGGATCACTGAGTCAAGAGA 2879
QY 141 TCGAGACCATCTGCGCAACATGTAAGAAACCCGCTTACTTAATAAATAATTAATGTC 200
DB 2878 TTGAGACCATCTGCGCAACATGTAAGAAACCCGCTTACTTAATAAATAATTAATGTC 2819
QY 201 TGGGATGATGCGACACACTGTAGTCCAGCTACTCAGAGCCGAGATTGAGTGAAGC 260
DB 2818 CAGGCTGTGTGTATGCTGCTGTATATCCAGCTACTCAGAGGCTGAGGTTGCAAGTGAAGC 2759
QY 261 TGAAGTGAAGTGAAGCCGAATCAAGATCAAGAGTGAAG 302
DB 2758 TGAAGTGAAGTGAAGCCGCTTCCAGCTGGGAGGAGAGTGAAG 2717

RESULT 38
US-11-266-748A-202571
Sequence 202571, Application US/11266748A
Publication No. US20060134663A1

GENERAL INFORMATION:
APPLICANT: Harkin, Paul
APPLICANT: Mulligan, Patrick
TITLE OF INVENTION: Transcription Microarray Technology and
TITLE OF INVENTION: Methods of Using the Same
FILE REFERENCE: 55815-0102 (319189)
CURRENT APPLICATION NUMBER: US/11/266,748A
CURRENT FILING DATE: 2005-11-03
PRIOR APPLICATION NUMBER: EP 04105479.2
PRIOR FILING DATE: 2004-11-03
PRIOR APPLICATION NUMBER: EP 04105482.6
PRIOR FILING DATE: 2004-11-03
PRIOR APPLICATION NUMBER: EP 04105483.4
PRIOR FILING DATE: 2004-11-03
PRIOR APPLICATION NUMBER: EP 04105507.0
PRIOR FILING DATE: 2004-11-03
PRIOR APPLICATION NUMBER: EP 04105485.9
PRIOR FILING DATE: 2004-11-03
PRIOR APPLICATION NUMBER: EP 04105484.2
PRIOR FILING DATE: 2004-11-03
PRIOR APPLICATION NUMBER: US 60/662,276
PRIOR FILING DATE: 2005-03-14

PRIOR APPLICATION NUMBER: US 60/700,293
PRIOR FILING DATE: 2005-07-18
NUMBER OF SEQ ID NOS: 48396
SOFTWARE: PatentIn version 3.3
SEQ ID NO 202571
LENGTH: 1000
TYPE: DNA
ORGANISM: Homo Sapiens
US-11-266-748A-202571

Query Match 35.4%; Score 141.6; DB 8; Length 1000;
Best Local Similarity 74.0%; Pred. No. 6.5e-21;
Matches 194; Conservative 1; Mismatches 60; Indels 7; Gaps 1;

QY 84 CTTGTAATCCAGCACTTGGGAGGCCAAGTGGCGGATCACTGAGTCAAGATCG 143
DB 383 CTTGTAATCCAGCACTTGGGAGGCCAAGTGGCGGATCACTGAGTCAAGATCG 442
QY 144 AGACATCTCGGCAACATGGTGAACCCCGCTTTTCTAATAAATPACAAAATAGCTCG 203
DB 443 AGACATCTCGGCAACATGGTGAACCCCGCTTTTCTAATAAATPACAAAATAGCTCG 502
QY 204 GCATGTGGCACAACCTGTAGTCCAGCTACTCAGAG-----CCGAGATTGAGT 256
DB 503 GCATGTGGCACAACCTGTAGTCCAGCTACTCAGAG-----CCGAGATTGAGT 562
QY 257 GAGCTGAGATCGCAGAGTGAAGCCCAATCACAATCAGAGTGAAGCAGACGCC 316
DB 563 GAACTTGGGAGGTGAAGTTGCAAGTGAAGCCAGCTCGGCTGGAGCAGAGTGAAGCTTT 622
QY 317 GTCTCAAAAACAACAACAAAAA 338
DB 623 GTCTCAAAAACAACAAAAA 644

RESULT 39

US-11-266-748A-23004/C
Sequence 23004, Application US/11266748A
Publication No. US20060134663A1
GENERAL INFORMATION:
APPLICANT: Harkin, Paul
APPLICANT: Johnston, Patrick
APPLICANT: Mulligan, Karl
TITLE OF INVENTION: Transcriptome Microarray Technology and
TITLE OF INVENTION: Methods of Using the Same
FILE REFERENCE: 55815-0102 (319189)
CURRENT APPLICATION NUMBER: US/11/266,748A
PRIOR APPLICATION NUMBER: EP 04105479.2
PRIOR FILING DATE: 2004-11-03
PRIOR APPLICATION NUMBER: EP 04105482.6
PRIOR FILING DATE: 2004-11-03
PRIOR APPLICATION NUMBER: EP 04105483.4
PRIOR FILING DATE: 2004-11-03
PRIOR APPLICATION NUMBER: EP 04105507.0
PRIOR FILING DATE: 2004-11-03
PRIOR APPLICATION NUMBER: EP 04105485.9
PRIOR FILING DATE: 2004-11-03
PRIOR APPLICATION NUMBER: EP 04105484.2
PRIOR FILING DATE: 2004-11-03
PRIOR APPLICATION NUMBER: US 60/662,276
PRIOR FILING DATE: 2005-03-14
PRIOR APPLICATION NUMBER: US 60/700,293
PRIOR FILING DATE: 2005-07-18
NUMBER OF SEQ ID NOS: 48396
SOFTWARE: PatentIn version 3.3
SEQ ID NO 23004
LENGTH: 216387
TYPE: DNA
ORGANISM: Homo Sapiens
US-11-266-748A-23004

Query Match 35.4%; Score 141.6; DB 8; Length 216387;

Best Local Similarity 69.1%; Pred. No. 8.5e-21;
Matches 192; Conservative 1; Mismatches 85; Indels 0; Gaps 0;

QY 66 CATGACACTGCTGATATGCTGTAATCCAGCACTTGGGAGGCCAAGTGGCGGATCA 125
DB 87903 CAAATAACAGGCGAGGCTGTAATCCAGCACTTGGGAGGCCAAGTGGCGGATCA 87844
QY 126 CTTGAGGTCAAGAGTTTGAACAGCTGGCCAAACATGTGAACCCCGTCTTACTAAA 185
DB 87843 CTTGAGGTCAAGAGTTTGAACAGCTGGCCAAACATGTGAACCCCGTCTTACTAAA 87784
QY 186 AATACAAAAATAGCTGGGAGTGGGCAACACCTGTAGTCCAGCTACTCAGAGCCG 245
DB 87783 AATACAAAAATAGCTGGGAGTGGGCAACACCTGTAGTCCAGCTACTCAGAGCCG 87724
QY 246 GAGATTGCAAGTGAAGTGAATCCAGAGTGAAGCCCAATCACAATCAGATCAAGATGAGCAG 305
DB 87723 ACCGCAAGAGAAATCACTGTAACCTGGGTGGAGGCTGCAATGAGCAGATCTGCCA 87664
QY 306 AGTGAGCKCCGCTCAAAAACAACAACAAAAA 343
DB 87663 CTATGGGTGAACAAAGGCTCAAAAACAAAAA 87626

RESULT 40

US-11-266-748A-201568
Sequence 201568, Application US/11266748A
Publication No. US20060134663A1
GENERAL INFORMATION:
APPLICANT: Harkin, Paul
APPLICANT: Johnston, Patrick
APPLICANT: Mulligan, Karl
TITLE OF INVENTION: Transcriptome Microarray Technology and
TITLE OF INVENTION: Methods of Using the Same
FILE REFERENCE: 55815-0102 (319189)
CURRENT APPLICATION NUMBER: US/11/266,748A
PRIOR APPLICATION NUMBER: EP 04105479.2
PRIOR FILING DATE: 2004-11-03
PRIOR APPLICATION NUMBER: EP 04105482.6
PRIOR FILING DATE: 2004-11-03
PRIOR APPLICATION NUMBER: EP 04105483.4
PRIOR FILING DATE: 2004-11-03
PRIOR APPLICATION NUMBER: EP 04105507.0
PRIOR FILING DATE: 2004-11-03
PRIOR APPLICATION NUMBER: EP 04105485.9
PRIOR FILING DATE: 2004-11-03
PRIOR APPLICATION NUMBER: EP 04105484.2
PRIOR FILING DATE: 2004-11-03
PRIOR APPLICATION NUMBER: US 60/662,276
PRIOR FILING DATE: 2005-03-14
PRIOR APPLICATION NUMBER: US 60/700,293
PRIOR FILING DATE: 2005-07-18
NUMBER OF SEQ ID NOS: 48396
SOFTWARE: PatentIn version 3.3
SEQ ID NO 201568
LENGTH: 632
TYPE: DNA
ORGANISM: Homo Sapiens
US-11-266-748A-201568

Query Match 35.4%; Score 141.4; DB 8; Length 632;
Best Local Similarity 77.1%; Pred. No. 7e-21;
Matches 172; Conservative 0; Mismatches 51; Indels 0; Gaps 0;

QY 26 CATGGAACCAAAATTAATAAGACATTGTGACGCGCAGCATGACATGGCTGAATGCC 85
DB 311 CAGGGAGGGAAGTTATGTGCAAAAAGTACAGGCCAGGCGCGGATGCTCAAGGCC 370
QY 86 TGTATCCAGCACTTGGGAGGCCAAGTGGCGGATCACTTGAAGTCAAGATGAG 145
DB 371 TATATCCAGCACTTGGGAGGCCAAGTGGCGGATCACTTGAAGTCAAGATGAG 430

QY 146 ACCATCTGGCCCAACATGGTGAACCCCTCTTACTAAATAATACAAAATAGCTGGCC 205
DB 431 ACCAGCTGGCCCAACATGGTGAACCCCTCTTACTAAATAATACAAAATAGCTGGCT 490
QY 206 ATGGTGGACACACCTGTATGCTCCAGCTACTCAGAGCCGAG 248
DB 491 ATGATGGCTCGGCTGTAGTCTCTAGTACTCAGAGGCTGAG 533

RESULT 41

US-11-266-748A-59943/c
Sequence 59943, Application US/11266748A
Publication No. US20060134663A1
GENERAL INFORMATION:

APPLICANT: Harlin, Paul
APPLICANT: Johnston, Patrick
TITLE OF INVENTION: Transcription Microarray Technology and
FILE REFERENCE: 55815-0102 (319189)
CURRENT APPLICATION NUMBER: US/11/266,748A
PRIOR FILING DATE: 2005-11-03
PRIOR APPLICATION NUMBER: EP 04105479.2
PRIOR FILING DATE: 2004-11-03
PRIOR APPLICATION NUMBER: EP 04105482.6
PRIOR FILING DATE: 2004-11-03
PRIOR APPLICATION NUMBER: EP 04105483.4
PRIOR FILING DATE: 2004-11-03
PRIOR APPLICATION NUMBER: EP 04105507.0
PRIOR FILING DATE: 2004-11-03
PRIOR APPLICATION NUMBER: EP 04105485.9
PRIOR FILING DATE: 2004-11-03
PRIOR APPLICATION NUMBER: EP 04105484.2
PRIOR FILING DATE: 2004-11-03
PRIOR APPLICATION NUMBER: US 60/662,276
PRIOR FILING DATE: 2005-03-14
PRIOR APPLICATION NUMBER: US 60/700,293
PRIOR FILING DATE: 2005-07-18
NUMBER OF SEQ ID NOS: 483996
SOFTWARE: Patent version 3.3
SEQ ID NO 59943
LENGTH: 2445
TYPE: DNA
ORGANISM: Homo Sapiens
US-11-266-748A-59943

Query Match 35.4%; Score 141.4; DB 8; Length 2445;
Best Local Similarity 88.0%; Pred. No. 7.5e-21;
Matches 154; Conservative 0; Mismatches 21; Indels 0; Gaps 0;

QY 81 ATGCTGTATCCAGCAGCTTCGGAGGCCCAAGGTGGCGGATCAGCTGAGGTCAAGGA 140
DB 344 AGGCTGTATCCAGCAGCTTCGGAGGCCCAAGGTGGCGGATCAGCTGAGGTCAAGGA 285
QY 141 TCGAGACCATCTGGCCCAACATGGTGAACCCCTCTTACTAAATAATACAAAATAGC 200
DB 284 TCGAGACCATCTGGCCCAACATGGTGAACCCCTCTTACTAAATAATACAAAATAGC 225
QY 201 TGGGATGTGGCACAACCTGTAGTCCAGCTACTCAGAGCCGAGATTGAG 255
DB 224 CAGGATGTGGCACAACCTGTAGTCCAGCTACTCAGAGCCGAGATTGAG 170

RESULT 42

US-10-517-441-32/c
Sequence 32, Application US/10517441
Publication No. US20060121467A1
GENERAL INFORMATION:

APPLICANT: FOEKENS, John
APPLICANT: HARBECK, Nedla
APPLICANT: KOENIG, Thomas
APPLICANT: MAIER, Sabine
APPLICANT: MARTENS, John

APPLICANT: MODEL, Fabian
APPLICANT: NIMMERICH, Inko
APPLICANT: RUJAN, Tamara
APPLICANT: SCHMITT, Armin
APPLICANT: SCHMITT, Manfred
APPLICANT: LOOK, Maxime P.
APPLICANT: MARX, Almut
APPLICANT: HOEFER, Heinz
TITLE OF INVENTION: Method and nucleic acids for the improved treatment of breast cel
FILE REFERENCE: 47675-93
CURRENT APPLICATION NUMBER: US/10/517,441
PRIOR FILING DATE: 2004-12-11
PRIOR APPLICATION NUMBER: PCT/EP2003/010881
PRIOR FILING DATE: 2003-10-01
PRIOR APPLICATION NUMBER: DE 10317955.0
PRIOR FILING DATE: 2003-04-17
PRIOR APPLICATION NUMBER: DE 10300096.8
PRIOR FILING DATE: 2003-01-07
PRIOR APPLICATION NUMBER: DE 10245779.4
PRIOR FILING DATE: 2002-10-01
NUMBER OF SEQ ID NOS: 2147
SEQ ID NO 32
LENGTH: 6521
TYPE: DNA
ORGANISM: Homo Sapiens
US-10-517-441-32

Query Match 35.4%; Score 141.4; DB 6; Length 6521;
Best Local Similarity 88.0%; Pred. No. 7.9e-21;
Matches 154; Conservative 0; Mismatches 21; Indels 0; Gaps 0;

QY 81 ATGCTGTATCCAGCAGCTTCGGAGGCCCAAGGTGGCGGATCAGCTGAGGTCAAGGA 140
DB 2279 ATGCTGTATCCAGCAGCTTCGGAGGCCCAAGGTGGCGGATCAGCTGAGGTCAAGGA 2220
QY 141 TCGAGACCATCTGGCCCAACATGGTGAACCCCTCTTACTAAATAATACAAAATAGC 200
DB 2219 TCGAGACCATCTGGCCCAACATGGTGAACCCCTCTTACTAAATAATACAAAATAGC 2160
QY 201 TGGGATGTGGCACAACCTGTAGTCCAGCTACTCAGAGCCGAGATTGAG 255
DB 2159 TGGGATGTGGCACAACCTGTAGTCCAGCTACTCAGAGCCGAGATTGAG 2105

RESULT 43

US-11-266-748A-60244
Sequence 60244, Application US/11266748A
Publication No. US20060134663A1
GENERAL INFORMATION:

APPLICANT: Harlin, Paul
APPLICANT: Johnston, Patrick
TITLE OF INVENTION: Transcription Microarray Technology and
FILE REFERENCE: 55815-0102 (319189)
CURRENT APPLICATION NUMBER: US/11/266,748A
PRIOR FILING DATE: 2005-11-03
PRIOR APPLICATION NUMBER: EP 04105479.2
PRIOR FILING DATE: 2004-11-03
PRIOR APPLICATION NUMBER: EP 04105482.6
PRIOR FILING DATE: 2004-11-03
PRIOR APPLICATION NUMBER: EP 04105483.4
PRIOR FILING DATE: 2004-11-03
PRIOR APPLICATION NUMBER: EP 04105507.0
PRIOR FILING DATE: 2004-11-03
PRIOR APPLICATION NUMBER: EP 04105485.9
PRIOR FILING DATE: 2004-11-03
PRIOR APPLICATION NUMBER: EP 04105484.2
PRIOR FILING DATE: 2004-11-03
PRIOR APPLICATION NUMBER: US 60/662,276
PRIOR FILING DATE: 2005-03-14
PRIOR APPLICATION NUMBER: US 60/700,293

```

; PRIOR FILING DATE: 2005-07-18
; NUMBER OF SEQ ID NOS: 483996
; SOFTWARE: PatentIn version 3.3
; SEQ ID NO: 60244
; LENGTH: 86654
; TYPE: DNA
; ORGANISM: Homo Sapiens
US-11-266-748A-60244

Query Match      35.4%; Score 141.4; DB 8; Length 86654;
Best Local Similarity 72.3%; Pred. No. 9e-21;
Matches 206; Conservative 1; Mismatches 57; Indels 21; Gaps 1;

QY 84 CCTGTAATCCAGACCTTCGGAGAGCCAGGTCGAGTCACTGAGGTGAAGATTCG 143
DB 22962 CCTGTAATCCAGACCTTCGGAGAGCCAGGTCGAGTCACTGAGGTGAAGATTCG 23021
QY 144 AGACCATCTGGCCCAACATGTTGAAACCCCGTCTTTACTAATAAATACAAAAATAGCTCG 203
DB 23022 AGACCATCTGGCCCAACATGTTGAAACCCCGTCTTTACTAATAAATACAAAAATAGCTCG 23081
QY 204 GCATGTGGCACAACCTGTATGTCCTCCAGCTACTAGAGCCGAGATTGCACTGAGCTGA 263
DB 23082 GCATGTGGCACAACCTGTATGTCCTCCAGCTACTAGAGCCGAGATTGCACTGAGCTGA 23141
QY 264 GATGCAAGATGAGCCGCAATCAAGATCAAGATGAGGCTGAGGAGAAATCACTT 303
DB 23142 GATGCAAGATGAGCCGCAATCAAGATCAAGATGAGGCTGAGGAGAAATCACTT 23201
QY 304 -AGAAGTGAAGACCCGCTCTCAAAAAACAACAACAAAAACAAAAA 347
DB 23202 AAGAGTGAAGACCCGCTCTCAAAAAACAACAACAAAAACAAAAA 23246

RESULT 44
US-11-266-748A-23291/c
; Sequence 23291, Application US/11266748A
; Publication No. US20060134663A1
; GENERAL INFORMATION:
; APPLICANT: Harkin, Paul
; APPLICANT: Johnston, Patrick
; APPLICANT: Mulligan, Karl
; TITLE OF INVENTION: Transcription Microarray Technology and
; FILE REFERENCE: 55815-0102 (319189)
; CURRENT FILING DATE: 2005-11-03
; PRIOR FILING DATE: 2004-11-03
; PRIOR APPLICATION NUMBER: EP 04105479.2
; PRIOR FILING DATE: 2004-11-03
; PRIOR APPLICATION NUMBER: EP 04105482.6
; PRIOR FILING DATE: 2004-11-03
; PRIOR APPLICATION NUMBER: EP 04105483.4
; PRIOR FILING DATE: 2004-11-03
; PRIOR APPLICATION NUMBER: EP 04105507.0
; PRIOR FILING DATE: 2004-11-03
; PRIOR APPLICATION NUMBER: EP 04105485.9
; PRIOR FILING DATE: 2004-11-03
; PRIOR APPLICATION NUMBER: EP 04105484.2
; PRIOR FILING DATE: 2004-11-03
; PRIOR APPLICATION NUMBER: US 60/662,276
; PRIOR FILING DATE: 2005-03-14
; PRIOR APPLICATION NUMBER: US 60/700,293
; PRIOR FILING DATE: 2005-07-18
; NUMBER OF SEQ ID NOS: 483996
; SOFTWARE: PatentIn version 3.3
; SEQ ID NO: 23291
; LENGTH: 113853
; TYPE: DNA
; ORGANISM: Homo Sapiens
US-11-266-748A-23291

Query Match      35.4%; Score 141.4; DB 8; Length 113853;
Best Local Similarity 88.0%; Pred. No. 9.1e-21;
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Matches 154; Conservative 0; Mismatches 21; Indels 0; Gaps 0;

QY 81 ATGCTTATATCCAGACCTTCGGAGAGCCAGGTCGAGTCACTGAGGTGAAGATTCG 140
DB 108547 ATGCTTATATCCAGACCTTCGGAGAGCCAGGTCGAGTCACTGAGGTGAAGATTCG 108488
QY 141 TCGAGACCATCTGGCCCAACATGTTGAAACCCCGTCTTTACTAATAAATACAAAAATAGC 200
DB 108487 TCGAGACCATCTGGCCCAACATGTTGAAACCCCGTCTTTACTAATAAATACAAAAATAGC 108428
QY 201 TGGGCATGTGGCACAACCTGTATGTCCTCCAGCTACTAGAGCCGAGATTGTCAG 255
DB 108427 TGGGCATGTGGCACAACCTGTATGTCCTCCAGCTACTAGAGCCGAGATTGTCAG 108373

RESULT 45
US-11-266-748A-58517/c
; Sequence 58517, Application US/11266748A
; Publication No. US20060134663A1
; GENERAL INFORMATION:
; APPLICANT: Harkin, Paul
; APPLICANT: Johnston, Patrick
; APPLICANT: Mulligan, Karl
; TITLE OF INVENTION: Transcription Microarray Technology and
; FILE REFERENCE: 55815-0102 (319189)
; CURRENT FILING DATE: 2005-11-03
; PRIOR FILING DATE: 2004-11-03
; PRIOR APPLICATION NUMBER: EP 04105479.2
; PRIOR FILING DATE: 2004-11-03
; PRIOR APPLICATION NUMBER: EP 04105482.6
; PRIOR FILING DATE: 2004-11-03
; PRIOR APPLICATION NUMBER: EP 04105483.4
; PRIOR FILING DATE: 2004-11-03
; PRIOR APPLICATION NUMBER: EP 04105507.0
; PRIOR FILING DATE: 2004-11-03
; PRIOR APPLICATION NUMBER: EP 04105485.9
; PRIOR FILING DATE: 2004-11-03
; PRIOR APPLICATION NUMBER: EP 04105484.2
; PRIOR FILING DATE: 2004-11-03
; PRIOR APPLICATION NUMBER: US 60/662,276
; PRIOR FILING DATE: 2005-03-14
; PRIOR APPLICATION NUMBER: US 60/700,293
; PRIOR FILING DATE: 2005-07-18
; NUMBER OF SEQ ID NOS: 483996
; SOFTWARE: PatentIn version 3.3
; SEQ ID NO: 58517
; LENGTH: 154394
; TYPE: DNA
; ORGANISM: Homo Sapiens
US-11-266-748A-58517

Query Match      35.4%; Score 141.4; DB 8; Length 154394;
Best Local Similarity 77.1%; Pred. No. 9.2e-21;
Matches 172; Conservative 0; Mismatches 51; Indels 0; Gaps 0;

QY 81 ATGCTTATATCCAGACCTTCGGAGAGCCAGGTCGAGTCACTGAGGTGAAGATTCG 140
DB 139360 ATGCTTATATCCAGACCTTCGGAGAGCCAGGTCGAGTCACTGAGGTGAAGATTCG 139301
QY 141 TCGAGACCATCTGGCCCAACATGTTGAAACCCCGTCTTTACTAATAAATACAAAAATAGC 200
DB 139301 TCGAGACCATCTGGCCCAACATGTTGAAACCCCGTCTTTACTAATAAATACAAAAATAGC 139241
QY 201 TGGGCATGTGGCACAACCTGTATGTCCTCCAGCTACTAGAGCCGAGATTGTCAGT 260
DB 139240 TGGGCATGTGGCACAACCTGTATGTCCTCCAGCTACTAGAGCCGAGATTGTCAGT 139181
QY 261 TGAAGATGAGGAGCCGCAATCAAGATCAAGATGAGGCTGAGGAGAAATCA 303
DB 139180 TGAAGATGAGGAGCCGCAATCAAGATCAAGATGAGGCTGAGGAGAAATCA 139138
```

RESULT 46
US-11-266-748A-22662/C
Sequence 22662, Application US/11266748A
Publication No. US20060134663A1
GENERAL INFORMATION:
APPLICANT: Harkin, Paul
APPLICANT: Johnston, Patrick
APPLICANT: Mulligan, Karl
TITLE OF INVENTION: Transcription Microarray Technology and
FILE REFERENCE: 55815-0102 (319189)
CURRENT FILING DATE: 2005-11-03
PRIOR APPLICATION NUMBER: EP 04105479.2
PRIOR FILING DATE: 2004-11-03
PRIOR APPLICATION NUMBER: EP 04105482.6
PRIOR FILING DATE: 2004-11-03
PRIOR APPLICATION NUMBER: EP 04105483.4
PRIOR FILING DATE: 2004-11-03
PRIOR APPLICATION NUMBER: EP 04105507.0
PRIOR FILING DATE: 2004-11-03
PRIOR APPLICATION NUMBER: EP 04105485.9
PRIOR FILING DATE: 2004-11-03
PRIOR APPLICATION NUMBER: EP 04105484.2
PRIOR FILING DATE: 2004-11-03
PRIOR APPLICATION NUMBER: US 60/662,276
PRIOR FILING DATE: 2005-03-14
PRIOR APPLICATION NUMBER: US 60/700,293
PRIOR FILING DATE: 2005-07-18
NUMBER OF SEQ ID NOS: 483996
SOFTWARE: PatentIn version 3.3
SEQ ID NO 22662
LENGTH: 164429
TYPE: DNA
ORGANISM: Homo Sapiens
US-11-266-748A-22662

Query Match 35.4%; Score 141.4; DB 8; Length 164429;
Best Local Similarity 77.1%; Pred. No. 9.3e-21;
Matches 172; Conservative 0; Mismatches 51; Indels 0; Gaps 0;

QY 81 ATGCGTGAATCCAGCACTTCGGAGGCGCAAGTGGCGGATCACTGAGTCAAGAGA 140
DB 27612 ATGCGTGAATCCAGCACTTCGGAGGCGTGAAGGCGGCAATCACTGAGTCAAGAGT 27553
QY 141 TCGAGACCACTCTGCGCAATGTGTAAACCCCGTCTTAAATAAATAAATAATAGC 200
DB 27552 TCGAGACCACTCTGCGCAATGTGTAAACCCCGTCTTAAATAAATAAATAATAGC 27493
QY 201 TGGGCAATGTGGGCAACACCTGTAGTCCAGCTACTCAAGAGCCGAGATTGCAGTAGC 260
DB 27492 CGGGCAATGTGGGCAACACCTGTATCCAGCTACTCAAGAGCCGAGATTGCAAGATTA 27433
QY 261 TGAGATCGACAGAGTGAAGCCGAATCAACAGATCAAGAGTAGC 303
DB 27432 CTGAACCCAGAGGAGGAGGTTCAGTGAACCAAGATCTGTC 27390

RESULT 47
US-11-266-748A-24937/C
Sequence 24937, Application US/11266748A
Publication No. US20060134663A1
GENERAL INFORMATION:
APPLICANT: Harkin, Paul
APPLICANT: Johnston, Patrick
APPLICANT: Mulligan, Karl
TITLE OF INVENTION: Transcription Microarray Technology and
FILE REFERENCE: 55815-0102 (319189)
CURRENT FILING DATE: 2005-11-03
PRIOR APPLICATION NUMBER: US/11/266,748A
PRIOR FILING DATE: 2004-11-03

PRIOR APPLICATION NUMBER: EP 04105482.6
PRIOR FILING DATE: 2004-11-03
PRIOR APPLICATION NUMBER: EP 04105483.4
PRIOR FILING DATE: 2004-11-03
PRIOR APPLICATION NUMBER: EP 04105507.0
PRIOR FILING DATE: 2004-11-03
PRIOR APPLICATION NUMBER: EP 04105485.9
PRIOR FILING DATE: 2004-11-03
PRIOR APPLICATION NUMBER: EP 04105484.2
PRIOR FILING DATE: 2004-11-03
PRIOR APPLICATION NUMBER: US 60/662,276
PRIOR FILING DATE: 2005-03-14
PRIOR APPLICATION NUMBER: US 60/700,293
PRIOR FILING DATE: 2005-07-18
NUMBER OF SEQ ID NOS: 483996
SOFTWARE: PatentIn version 3.3
SEQ ID NO 24937
LENGTH: 86215
TYPE: DNA
ORGANISM: Homo Sapiens
US-11-266-748A-24937

Query Match 35.3%; Score 141.2; DB 8; Length 86215;
Best Local Similarity 67.9%; Pred. No. 9.9e-21;
Matches 197; Conservative 0; Mismatches 93; Indels 0; Gaps 0;

QY 14 CATGTCTGGGCAATGGAAACCAATTTAATAAGACATTGTCAAGCCAGGATGAC 73
DB 67383 CATTTATTTCTTTTAAATGAACAAATTTAGAGAAAGTTATTCAGGCCAGGAT 67324
QY 74 TGGCGAATGCTGTATCCAGCACTTCGGAGGCGCAAGTGGCGGATCACTGAGGT 133
DB 67323 GGTATCAATGCTGTATCCAGCACTTTAGAGGCTGAGCGGGTGTATCACTGAGGT 67284
QY 134 CAAGAGATGAGACCATCTGCGCAATGCTGAACCCCGTCTTAAATAAATAACAA 193
DB 67263 CAGAGATTCGAGACCAAGCTTACCAATGCTGCAACCTGTCTTAAATAAATAACAA 67204
QY 194 AATATGCTGGGCAATGTGGGCAACACCTGTATGCTCCAGCTACTCAAGAGCCGAGATTGC 253
DB 67203 AATATGCTGGGCAATGTATGATGTAGTCTGTATGCTCCAGCTACTCAAGAGGCTGAGGTAG 67144
QY 254 AGTAGCTGAGATGCGAGAGTGAACCGAAATCAAGATCAAGAGTAGC 303
DB 67143 ACAATCACTGAACCCAGAGGCGAGGTTGCACTGAGTGAATGTGC 67094

RESULT 48
US-11-266-748A-119261
Sequence 119261, Application US/11266748A
Publication No. US20060134663A1
GENERAL INFORMATION:
APPLICANT: Harkin, Paul
APPLICANT: Johnston, Patrick
APPLICANT: Mulligan, Karl
TITLE OF INVENTION: Transcription Microarray Technology and
FILE REFERENCE: 55815-0102 (319189)
CURRENT FILING DATE: 2005-11-03
PRIOR APPLICATION NUMBER: EP 04105479.2
PRIOR FILING DATE: 2004-11-03
PRIOR APPLICATION NUMBER: EP 04105482.6
PRIOR FILING DATE: 2004-11-03
PRIOR APPLICATION NUMBER: EP 04105483.4
PRIOR FILING DATE: 2004-11-03
PRIOR APPLICATION NUMBER: EP 04105507.0
PRIOR FILING DATE: 2004-11-03
PRIOR APPLICATION NUMBER: EP 04105485.9
PRIOR FILING DATE: 2004-11-03
PRIOR APPLICATION NUMBER: US/11/266,748A
PRIOR FILING DATE: 2004-11-03
PRIOR APPLICATION NUMBER: EP 04105484.2
PRIOR FILING DATE: 2004-11-03
PRIOR APPLICATION NUMBER: US 60/662,276

```

: PRIOR FILING DATE: 2005-03-14
: PRIOR APPLICATION NUMBER: US 60/700,293
: NUMBER OF SEQ ID NOS: 48396
: SOFTWARE: PatentIn version 3.3
: SEQ ID NO 119261
: LENGTH: 1000
: TYPE: DNA
: ORGANISM: Homo Sapiens
US-11-266-748A-119261

Query Match      35.3%; Score 141; DB 8; Length 1000;
Best Local Similarity 68.8%; Pred. No. 8.7e-21;
Matches 229; Conservative 1; Mismatches 86; Indels 17; Gaps 2;

QY 81 ATGCTGTAAATCCAGACCTTCGGAGGCGCAAGTGGGCGGATCACTGAGTCAAGAGA 140
DB 200 ACGCTGTAAATCCAGACCTTCGGAGGCGCGCTGGAGGCGGATTCCTGAGGTGAGAGT 259
QY 141 TCGAGACCATCTGCGCAACATGTGGAAACCCCGCTTACTAAATAACAAATAATAGC 200
DB 260 TCGAGACCATCTGCGCAACATGTGGAAACCCCGCTTACTAAATAACAAATAATAGC 319
QY 201 TGGGCGATGTGGACACACCTGTAGTCCAGACTACTACAGA-GCCGAGATTGCAGTGAG 259
DB 320 TGGGCGATGTGGACACACCTGTAGTCCAGACTCTCGGAGGCGAGGAATCGCTTGA 379
QY 260 CTGAGATCGCAGAGTGAAGCCGAATCACAATCACA-----GAGTGAGC 303
DB 380 CCGGAGGTAGAGGTTCAGATGAGCCGAGATCGCACTACCTCCAGCTGGGCGGAC 439
QY 304 AGAGTGAGACKCCGCTCTCAAAAACAACAACAAACAAACAAATTAAGCATTTGCC 363
DB 440 AGAGCAAGACTCTCTTCAAAAAAAGGCGGCGGCGGTGCT 499
QY 364 ATCTGCGGTTCCTCCAGACTATTGACGAGACCAA 396
DB 500 CACTCTGTAAATCCAGACCTCTGAAGCGCA 532

RESULT 49
US-11-266-748A-161425/c
: Sequence 161425, Application US/11266748A
: Publication No. US20060134663A1
: GENERAL INFORMATION:
: APPLICANT: Johnston, Patrick
: APPLICANT: Mulligan, Karl
: TITLE OF INVENTION: Transcriptome Microarray Technology and
: FILE REFERENCE: 55815-0102 (319189)
: CURRENT FILING DATE: US/11/266,748A
: PRIOR APPLICATION NUMBER: EP 04105479.2
: PRIOR FILING DATE: 2004-11-03
: PRIOR APPLICATION NUMBER: EP 04105482.6
: PRIOR FILING DATE: 2004-11-03
: PRIOR APPLICATION NUMBER: EP 04105483.4
: PRIOR FILING DATE: 2004-11-03
: PRIOR APPLICATION NUMBER: EP 04105507.0
: PRIOR FILING DATE: 2004-11-03
: PRIOR APPLICATION NUMBER: EP 04105485.9
: PRIOR FILING DATE: 2004-11-03
: PRIOR APPLICATION NUMBER: EP 04105484.2
: PRIOR FILING DATE: 2004-11-03
: PRIOR APPLICATION NUMBER: US 60/662,276
: PRIOR FILING DATE: 2005-03-14
: PRIOR APPLICATION NUMBER: US 60/700,293
: PRIOR FILING DATE: 2005-07-18
: NUMBER OF SEQ ID NOS: 48396
: SOFTWARE: PatentIn version 3.3
: SEQ ID NO 161425
: LENGTH: 1000
```

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: TYPE: DNA
: ORGANISM: Homo Sapiens
US-11-266-748A-161425

Query Match      35.3%; Score 141; DB 8; Length 1000;
Best Local Similarity 68.8%; Pred. No. 8.7e-21;
Matches 229; Conservative 1; Mismatches 86; Indels 17; Gaps 2;

QY 81 ATGCTGTAAATCCAGACCTTCGGAGGCGCAAGTGGGCGGATCACTGAGTCAAGAGA 140
DB 801 ACGCTGTAAATCCAGACCTTCGGAGGCGCGCTGGAGGCGGATTCCTGAGGTGAGAGT 742
QY 141 TCGAGACCATCTGCGCAACATGTGGAAACCCCGCTTACTAAATAACAAATAATAGC 200
DB 741 TCGAGACCATCTGCGCAACATGTGGAAACCCCGCTTACTAAATAACAAATAATAGC 682
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RESULT 50
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: LENGTH: 1000
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: ORGANISM: Homo Sapiens
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Best Local Similarity 68.8%; Pred. No. 8.7e-21;
Matches 229; Conservative 1; Mismatches 86; Indels 17; Gaps 2;
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OY 81 ATGCTGTATATCCAGACATTGCGAGGCGAAGGTGGGGGATCACTGAGTCAAGAGA 140
DB 200 AGGCTGTATATCCAGACATTGCGAGGCGCTGGAGGGTGGATTGCTTGAGTCAAGAGT 259
OY 141 TCGAGACCATCTGGCCAAATGTGTAAACCCCGTCTTTACTTAAAAATACAAAAATAGC 200
DB 260 TCGAGACGAGCCTGCTTAATGTGTAAACCCCGTCTCACTTAAAAATACAAAAATAGC 319
OY 201 TGGGCAATGGTGGCACACACTGTAGTCCCACTACTCAGGA-GCCGAGATTGCAGTAG 259
DB 320 TGGGCGTGGTGGCACACGCTGTAGTCCCACTCTCGGGAGGCAAGAGATGCTTGAA 379
OY 260 CTGAGATGCGAGGTGAGCGGAAATGACAGATGCA-----GAGTGAGC 303
DB 380 CCCCAGAGGTAGAGGTTGCAGTGAGCGAGATGCACTACACTCCAGCCTGGGCGAC 439
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DB 500 CACTCTGTATCCAGACACTAGAGGCCGA 532

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